

Applicants : David M. Stern, et al.
U.S. Serial No: 09/872,185
Filed : June 1, 2001
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REMARKS

Claims 1-3 and 7-18 are pending and claims 3, 7-13 and 16-18 are under examination. Claim 3 has been amended. Applicants maintain that this amendment raises no issue of new matter and is fully supported by the specification as filed. Applicants respectfully request that this Amendment be entered. Upon entry of this Amendment, claims 1-3 and 7-18 will still be pending and claims 3, 7-13 and 16-18 will still be under examination.

Rejection under 35 U.S.C. §103(a)

The Examiner rejected claims 3-11 under 35 U.S.C. §103(a) as allegedly obvious over United States Patent No. 5,864,018 ("Morser") in view of Ritthauer et al. (1995) ("Ritthauer").

Specifically, the Examiner asserts that it would have been prima facie obvious at the time of the invention to treat inflammation in a subject by inhibiting the interaction of AGE and RAGE by administering sRAGE or the anti-RAGE antibodies disclosed by Morser, because Morser demonstrates that sRAGE or anti-RAGE antibodies block the interaction between RAGE and its ligands and Ritthauer teaches that RAGE and its ligands are involved in the diseases involving inflammation.

In response to the Examiner's rejection of claims 4-6, but without conceding the correctness thereof, applicants point out that these claims were canceled in the September 17, 2003 Amendment in Response to June 17, 2003 Office Action, thereby rendering the

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rejection thereof moot.

Further, applicants respectfully traverse the rejection of claims 3 and 7-11. In order to find the subject application obvious over Morser in view of Ritthauer, the prior art references, in combination, must teach or suggest all the elements thereof, and create both a motive to combine and a reasonable expectation of success. Morser and Ritthauer fail to do this.

Claims 3 and 7-11, as amended, provide a method for treating inflammation in a subject which comprises administering to the subject an agent selected from the group consisting of soluble receptor for advanced glycation endproduct (sRAGE) or anti-RAGE antibody or anti-EN-RAGE F(ab')₂ fragment in an amount which inhibits the interaction between receptor for advanced glycation endproduct (RAGE) and either AGE or EN-RAGE, thereby treating inflammation in the subject.

This invention is based on the surprising discovery that inhibiting the interaction of RAGE and either AGE or EN-RAGE is useful in treating inflammation. Prior to applicants' surprising discovery, *as evidenced by the experimental data in this application*, there would have been no reasonable expectation that inhibiting the interaction of RAGE and either AGE or EN-RAGE would succeed in treating inflammation. The Examiner's assertion of obviousness ignores this point.

The Examiner has in no way shown how Morser in view of Ritthauer teach or suggest applicants' invention which is based on this surprising finding. To maintain otherwise would constitute

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hindsight.

Accordingly, applicants maintain that the subject claims are not obvious over Morser in view Ritthauer and therefore satisfy the requirements of 35 U.S.C. §103(a).

Further, the Examiner rejected claims 12, 13 and 16-18 under 35 U.S.C. §103(a) as allegedly obvious over Morser and Ritthauer in view of United States Patent No. 5,998,408 ("Baker").

Specifically, the Examiner asserts that it would have been prima facie obvious at the time of the invention to treat inflammation associated with autoimmune disease in a subject by inhibiting the interaction of AGE and RAGE by administering sRAGE or the anti-RAGE antibodies disclosed by Morser. This allegedly is because Morser demonstrates that sRAGE or anti-RAGE antibodies block the interaction between RAGE and its ligands, Ritthauer teaches that RAGE and its ligands are involved in the diseases involving inflammation and Baker teaches diseases involving inflammation, specifically the autoimmune process.

In response to the Examiner's rejection of claims 12, 13 and 16-18, applicants respectfully traverse the rejection of these claims.

Claims 12, 13 and 16-18 are discussed above.

The non-obviousness of the rejected claims over Morser and Ritthauer is discussed above. Baker, which according to the

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Examiner merely teaches diseases involving inflammation, does not cure the deficiency of the other two references, in that it does not teach applicants' surprising discovery underlying the claimed invention.

Accordingly, applicants maintain that the subject claims are not obvious over Morser and Ritthauer in view of Baker, and therefore satisfy the requirements of 35 U.S.C. §103(a).

Obviousness-Type Double Patenting Rejection

The Examiner provisionally rejected claims 3-11 and 16 as allegedly unpatentable under the judicially created doctrine of obviousness-type double patenting over claims 47, 50, 55-60, 62-65 and 67 of copending U.S. Application No. 09/167,705 in view of United States Patent No. 5,864,018 and Ritthauer et al. (1995). According to the Examiner, a timely filed terminal disclaimer in compliance with 37 C.F.R. §1.321(c) may be used to overcome a provisional rejection based on a nonstatutory double patenting ground.

In response, but without conceding the correctness of the Examiner's rejection, applicants will consider submitting a terminal disclaimer for claims 3-11 and 16 once the rejection is no longer provisional.

Supplemental Information Disclosure Statement

This Information Disclosure Statement is submitted as a supplement to the Information Disclosure Statement filed March 29, 2002.

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In accordance with their duty of disclosure under 37 C.F.R. §1.56, applicants would like to direct the Examiner's attention to U.S. Provisional Application No. 60/469,428, filed May 9, 2003 (**Exhibit 1**) and United States Patent Application No. 10/840,927, filed May 7, 2004 (**Exhibit 2**). These references are listed on the attached Form PTO-1449 (**Exhibit A**).

Summary

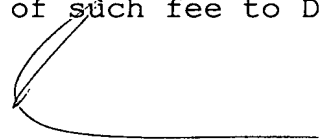
For the reasons set forth hereinabove, applicants respectfully request that all the claims of this application be allowed, and that the application proceed to issuance.

If a telephone interview would be of assistance in advancing prosecution of the subject application, applicants' undersigned attorneys invite the Examiner to telephone them at the number provided below.

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If a telephone interview would be of assistance in advancing prosecution of the subject application, applicants' undersigned attorneys invite the Examiner to telephone them at the number provided below.

No fee, other than the enclosed RCE filing fee, is deemed necessary in connection with the filing of this Preliminary Amendment and Supplemental Information Disclosure Statement. However, if any additional fee is required, authorization is hereby given to charge the amount of such fee to Deposit Account No. 03-3125.



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Form PTO-1449 U.S. Department of Commerce Patent and Trademark Office INFORMATION DISCLOSURE CITATION (Use several sheets if necessary)		Atty. Docket No. 575-64080 JPW/AJM/AAB		Serial No. 09/872,185	
		Applicant(s) David M. Stern, et al.			
		Filing Date June 1, 2004		Group Art Unit 1647	

U.S. PATENT DOCUMENTS									
Examiner Initials	Document Number	Date	Name	Class	Sub-class	Filing Date If Appropriate			
	6 0 4 6 9 4 2 8	05/09/2003	Ann Marie Schmidt, et al.						
	1 0 8 4 0 9 2 7	05/07/2004	Ann Marie Schmidt, et al.						

FOREIGN PATENT DOCUMENTS									
		Document Number	Date	Country	Class	Subclass	Translation		
							Yes	No	

OTHER DOCUMENTS (Including Author, Title, Date, Pertinent Pages, Etc.)		

EXAMINER	DATE CONSIDERED
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*EXAMINER: Initial if citation considered, whether or not citation is in conformance with MPEP 609: Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

*Provisional Application
for
United States Letters Patent*

To all whom it may concern:

Be it known that

Ann Marie Schmidt and David M. Stern

have invented certain new and useful improvements in

**RAGE AND RAGE G82S-RELATED METHODS
AND COMPOSITIONS FOR TREATING
INFLAMMATORY DISORDERS**

of which the following is a full, clear and exact description.

RAGE AND RAGE G82S-RELATED METHODS AND COMPOSITIONS
FOR TREATING INFLAMMATORY DISORDERS

5

Throughout this application, various publications are referenced. Full citations for these publications may be found immediately preceding the claims. The disclosures of these publications are hereby
10 incorporated by reference into this application in order to more fully describe the state of the art as of the date of the invention described and claimed herein.

Background of the Invention

15

The Receptor for Advanced Glycation Endproducts (RAGE), a multi-ligand member of the immunoglobulin superfamily of cell surface molecules, has been implicated in amplification of proinflammatory responses (1).
20 S100/calgranulins, proinflammatory signal transduction ligands of the receptor, are enriched in joints of subjects with rheumatoid arthritis ("RA") (2-4). Members of this family of molecules, long-associated with classic immune/inflammatory disorders (5-6), may be
25 released by activated inflammatory effector cells such as monocytes (7), thereby freeing them to engage cell surface RAGE and amplify host inflammatory responses. Indeed, accumulation of S100/calgranulins in synovial fluid and plasma of subjects with RA has been
30 correlated with indices of disease severity, such as bony erosions (4).

Recent studies have highlighted the possibility that polymorphisms within key domains of RAGE may influence
35 its function, so that under conditions of increased

ligand accumulation, individuals may be predisposed to heightened inflammatory responses (8-9). Importantly, a polymorphism of the RAGE gene has been identified within the V-type immunoglobulin domain in the extracellular region of the receptor, consisting of a glycine to serine change at position 82 (8). In previous studies, it was found that the ligands of RAGE, including S100/calgranulins, engage the V-domain of the receptor and activate signal transduction pathways, thereby modulating gene expression (1,10-14).

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Summary of the Invention

This invention provides a method for inhibiting binding between RAGE and a ligand thereof comprising contacting
5 the RAGE with an agent selected from the group consisting of soluble RAGE, soluble RAGE G82S, and an antibody directed to RAGE.

This invention further provides a method for inhibiting
10 binding between RAGE G82S and a ligand thereof comprising contacting the RAGE G82S with an agent selected from the group consisting of soluble RAGE, soluble RAGE G82S, and an antibody directed to RAGE G82S.

15 This invention further provides a method for treating an inflammatory disorder in a subject comprising administering to the subject a therapeutically effective amount of an agent that inhibits binding
20 between RAGE and/or RAGE G82S and a ligand thereof.

This invention further provides a method for inhibiting the onset of an inflammatory disorder in a subject comprising administering to the subject a
25 prophylactically effective amount of an agent that inhibits binding between RAGE and/or RAGE G82S and a ligand thereof.

This invention further provides a composition
30 comprising a pharmaceutically acceptable carrier and soluble RAGE G82S.

This invention further provides a method for inhibiting the onset of an inflammatory disorder in a subject

comprising administering to the subject a prophylactically effective amount of a nucleic acid which specifically inhibits the expression of RAGE or RAGE G82S in the subject's cells expressing same.

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This invention further provides a method for treating a subject afflicted with an inflammatory disorder comprising administering to the subject a therapeutically effective amount of a nucleic acid
10 which specifically inhibits the expression of RAGE or RAGE G82S in the subject's cells expressing same.

Finally, this invention provides an article of manufacture comprising a packaging material having
15 therein an agent that inhibits binding between RAGE and/or RAGE G82S and a ligand thereof, wherein the packaging material has affixed thereto a label indicating a use for the agent for treating an inflammatory disorder in a subject.

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Brief Description of the Figures

Figures 1A-1N. Induction of arthritis by bovine type II collagen in DBA/1 mice: increased expression of RAGE and S100/calgranulins. Six weeks after immunization (day 42), joint tissue from the hind feet (a-l) or stifle joint (m-n) was prepared for study.

1A-1L. Histology. In a-b-c, tissue was subjected to H&E analysis. Immunohistochemistry using anti-RAGE IgG (d-e-f); anti-S100/calgranulin IgG (g-h-i); or rabbit nonimmune IgG (j-k-l) was performed. Scale bar: a-b-c, 300 μ m; and d-l, 60 μ m.

1M-1N. Immunoblotting. Lysates were prepared from stifle joints and subjected to immunoblotting using anti-RAGE IgG (m); or anti-S100/calgranulin IgG (n). N=5 mice per condition were employed. Densitometric analysis of band intensity from all mice in each group was performed, and the mean \pm SD is shown. Representative bands are shown. Statistical considerations are indicated in the figure.

Figures 2A-2D. Blockade of RAGE suppresses inflammation in DBA/1 mice immunized/challenged with bovine type II collagen.

2A. Clinical score. At the indicated times after immunization with bovine type II collagen and treatment with murine sRAGE or vehicle (murine serum albumin (MSA) or PBS), clinical scoring was performed by blinded observers. The mean \pm standard deviation (SD) is shown; n= 10 mice per group. * indicates $p < 0.05$.

2B. Histologic score. Histologic scoring of joints in four limbs was performed on H&E-stained sections from mice six weeks after immunization. The mean \pm SD is shown; n=10 mice per group. * indicates $p < 0.05$.

5

2C. Ear swelling. Six weeks after immunization with bovine type II collagen, murine serum albumin (MSA)-, PBS- and sRAGE-treated mice were injected with bovine type II collagen into ear tissue. Ear thickness was
10 measured by a blinded observer immediately prior to local injection, and 18 hrs later. The mean \pm SD is shown; n=5 mice per group. * indicates $p < 0.05$ vs ear thickness in PBS- and MSA-treated groups.

15 2D. Antibody response to bovine type II collagen. At the indicated time points after immunization, plasma was retrieved from DBA/1 mice and subjected to ELISA for determination of total serum IgG levels to type II collagen. There were no statistically-significant
20 differences in total IgG levels among the groups.

Figures 3A-3F. Blockade of RAGE suppresses generation of cytokines and MMPs in DBA/1 mice immunized/challenged with bovine type II collagen.

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3A-3B. Joint tissue. Stifle joint tissue was retrieved six weeks after immunization with bovine type II collagen. Lysates were subjected to ELISA for detection of murine TNF-alpha (a) and IL-6 (b).
30 Results were normalized per mg/tissue extract.

3C-3D. Plasma. Plasma from mice was subjected to ELISA for levels of TNF-alpha (c) and IL-6 (d). In

a-b-c-d, n=10 mice per group. The mean \pm SD is shown. In a-d, * indicates $p < 0.05$.

3E-3F. Zymography. Stifle joint tissue was retrieved
5 and lysates were prepared and zymography for detection
of activity of MMP-2, -9 or -13 was performed. N=10
mice/group. Densitometric analysis of band intensity
from all mice/groups was performed, and the mean \pm SD
is shown. Representative bands are shown. In f, *
10 indicates $p < 0.05$ vs MSA-treated mice with collagen-
induced arthritis.

Figures 4A-4F. CHO cells bearing RAGE 82S display
increased affinity and cellular responsiveness to EN-
15 RAGE. CHO cells, which endogenously do not express
RAGE antigen, were stably-transfected with pcDNA3.1
vector containing cDNA encoding human RAGE 82G or the
variant allele 82S.

20 4A. Immunoblotting. Lysates of stably-transfected CHO
cells were prepared and subjected to immunoblotting
using anti-human RAGE IgG.

4B-4C. Radioligand binding assays. Purified EN-RAGE
25 was radiolabelled using 125 I and radioligand binding
assays were performed in 96-well tissue culture dishes
containing the indicated transfected CHO cells.
Equilibrium binding data were analyzed according to the
equation of Klotz and Hunston. Where indicated,
30 pretreatment with either antibodies, human soluble RAGE
or bovine serum albumin was performed. The mean \pm
standard deviation (SD) is shown. In c, * indicates
 $p < 0.01$ versus respective controls.

4D-4E. Activation of MEK 1/2 and p44/p42 MAP kinases. The indicated stably-transfected CHO cells were incubated with EN-RAGE for one hr. Immunoblotting of cell lysates was performed using anti-phosphorylated
5 MEK 1/2 (d) or p44/p42 MAP kinase (e). Where indicated, pretreatment with either BSA or sRAGE, or the indicated IgG for 2 hrs was performed. Control immunoblotting using antibody to total MEK 1/2 and p44/p42 MAP kinase revealed that there were no
10 differences in levels of total MEK 1/2 or p44/p42 (not shown).

4F. Activation of NF- κ B. Nuclear extracts were prepared from the indicated stably-transfected CHO
15 cells incubated with EN-RAGE for 6 hrs and EMSA was performed. Where indicated, cells were pretreated with either nonimmune/anti-RAGE IgG, soluble RAGE or BSA for 2 hrs prior to incubation with EN-RAGE. In d-e-f, bands were scanned into a densitometer, and band
20 density was quantified using ImageQuant. These experiments were performed at least three times and subjected to statistical analyses as indicated in the text and the figure legend. Representative experiments are shown.

25

Figures 5A-5H. Human peripheral blood mononuclear phagocytes (MPs) expressing RAGE 82S display increased responsiveness to EN-RAGE. MPs were purified from the blood of human subjects bearing RAGE (G82G) (n=5) or
30 heterozygotes (G82S) (n=5).

5A-5D. Activation of MEK 1/2 and p44/p42 MAP kinases. The indicated MPs were incubated with EN-RAGE or no mediator for one hr. Cell lysates were prepared and

immunoblotting performed using anti-phosphorylated MEK 1/2 (a,b) or p44/p42 MAP kinase (c,d). Densitometric analysis was performed and mean \pm SD is shown in b and d. Immunoblotting using antibody to total MEK 1/2 and p44/p42 MAP kinase revealed that there were no differences in levels of total MEK 1/2 or p44/p42 (not shown).

5E-5F. Generation of TNF-alpha and IL-6. Human MPs bearing the indicated RAGE alleles were cultured in the presence of either no mediator or EN-RAGE for 14 hrs. Supernatants were retrieved and levels of TNF-alpha (e) and IL-6 (f) determined by ELISA. The mean \pm SD is shown.

5G-5H. MMP-9 activity. Human MPs bearing the indicated RAGE alleles were cultured in the presence of either no mediator or EN-RAGE for 14 hrs. Supernatants were retrieved and subjected to zymography to assess levels of activated MMP-9. In b, d, and h, bands from n=5 G82G and n=5 G82S subjects were scanned into a densitometer and band density was quantified. Representative experiments are shown. The mean \pm SD is shown. In b,d,e,f and h, * indicates $p < 0.01$ versus baseline.

Detailed Description of the Invention

Definitions

5 "Antibody" shall include, by way of example, both naturally occurring and non-naturally occurring antibodies. Specifically, this term includes polyclonal and monoclonal antibodies, and antigen-binding fragments thereof. Furthermore, this term includes chimeric
10 antibodies and wholly synthetic antibodies, and antigen-binding fragments thereof.

"Anti-sense nucleic acid" shall mean any nucleic acid which, when introduced into a cell, specifically
15 hybridizes to at least a portion of an mRNA in the cell encoding a protein ("target protein") whose expression is to be inhibited, and thereby inhibits the target protein's expression.

20 "Catalytic nucleic acid" shall mean a nucleic acid that specifically recognizes a distinct substrate and catalyzes the chemical modification of this substrate. Catalytic nucleic acids include, for example, DNAzymes and ribozymes.

25

"DNAzyme" shall mean a catalytic nucleic acid that is DNA or whose catalytic component is DNA, and which specifically recognizes and cleaves a distinct target nucleic acid sequence, which can be either DNA or RNA.
30 Each DNAzyme has a catalytic component (also referred to as a "catalytic domain") and a target sequence-binding component consisting of two binding domains, one on either side of the catalytic domain.

"Expressible nucleic acid" shall mean a nucleic acid encoding a nucleic acid of interest and/or a protein of interest, which nucleic acid is an expression vector, plasmid or other construct which, when placed in a cell, permits the expression of the nucleic acid or protein of interest. Expression vectors and plasmids are well known in the art.

As used herein, "inflammatory disorder" includes, without limitation, any disorder possessing an inflammatory component. Examples of inflammatory disorders include autoimmune disorders generally, rheumatoid arthritis, amyotrophic lateral sclerosis, vascular disorders (e.g. atherosclerosis or transplant vasculopathy), and diabetes.

As used herein, "inhibit," when used in connection with the binding between RAGE and/or RAGE G82S with a ligand thereof, shall mean to reduce such binding. In one embodiment, "inhibit" shall mean to eliminate such binding.

"Inhibiting" the onset of a disorder shall mean either lessening the likelihood of the disorder's onset, or preventing the onset of the disorder entirely. In the preferred embodiment, inhibiting the onset of a disorder means preventing its onset entirely.

"Nucleic acid" shall mean any nucleic acid molecule, including, without limitation, DNA, RNA and hybrids thereof. The nucleic acid bases that form nucleic acid molecules can be the bases A, C, G, T and U, as well as derivatives thereof. Derivatives of these bases are well known in the art, and are exemplified in PCR

Systems, Reagents and Consumables (Perkin Elmer Catalogue 1996-1997, Roche Molecular Systems, Inc., Branchburg, New Jersey, USA).

5 "Ribozyme" shall mean a catalytic nucleic acid molecule which is RNA or whose catalytic component is RNA, and which specifically recognizes and cleaves a distinct target nucleic acid sequence, which can be either DNA or RNA. Each ribozyme has a catalytic component (also
10 referred to as a "catalytic domain") and a target sequence-binding component consisting of two binding domains, one on either side of the catalytic domain.

"Specifically hybridize" to a nucleic acid shall mean,
15 with respect to a first nucleic acid, that the first nucleic acid hybridizes to a second nucleic acid with greater affinity than to any other nucleic acid.

"Specifically inhibit" the expression of a protein
20 shall mean to inhibit that protein's expression (a) more than the expression of any other protein, or (b) more than the expression of all but 10 or fewer other proteins.

25 "Subject" shall mean any animal, such as a human, non-human primate, mouse, rat, guinea pig or rabbit.

"Treating" a disorder shall mean slowing, stopping or reversing the disorder's progression. In the preferred
30 embodiment, treating a disorder means reversing the disorder's progression, ideally to the point of eliminating the disorder itself. As used herein,

ameliorating a disorder and treating a disorder are equivalent.

Embodiments of the Invention

5

This invention provides methods for inhibiting binding between RAGE and/or RAGE G82S with a ligand thereof, and for inhibiting the onset of and treating inflammatory disorders such as rheumatoid arthritis.

10 This invention is based on the surprising discovery of a correlation between suppressing rheumatoid arthritis and blocking RAGE and/or RAGE G82S function.

Specifically, this invention provides a method for
15 inhibiting binding between RAGE and a ligand thereof comprising contacting the RAGE with an agent selected from the group consisting of soluble RAGE, soluble RAGE G82S, and an antibody directed to RAGE. This invention also provides a method for inhibiting binding between
20 RAGE G82S and a ligand thereof comprising contacting the RAGE G82S with an agent selected from the group consisting of soluble RAGE, soluble RAGE G82S, and an antibody directed to RAGE G82S.

25 In one embodiment of the instant methods, the agent is soluble RAGE. In another embodiment, the agent is soluble RAGE G82S. In a further embodiment, the agent is an antibody directed to RAGE. In yet a further embodiment, the agent is an antibody directed to RAGE
30 G82S, or a small molecule such as an organic molecule.

This invention also provides a method for treating an inflammatory disorder in a subject comprising administering to the subject a therapeutically

effective amount of an agent that inhibits binding between RAGE and/or RAGE G82S and a ligand thereof. This invention also provides a method for inhibiting the onset of an inflammatory disorder in a subject

5 comprising administering to the subject a prophylactically effective amount of an agent that inhibits binding between RAGE and/or RAGE G82S and a ligand thereof.

10 In the preferred embodiment of the instant methods, the subject is human. In one embodiment the inflammatory disorder is rheumatoid arthritis. In one embodiment of the instant methods, the agent is soluble RAGE. In another embodiment, the agent is soluble RAGE G82S. In

15 a further embodiment, the agent is an antibody directed to RAGE. In yet a further embodiment, the agent is an antibody directed to RAGE G82S, or a small molecule such as an organic molecule.

20 Determining a therapeutically or prophylactically effective amount of agent can be done based on animal data using routine computational methods. In one embodiment, the therapeutically or prophylactically effective amount contains between about 1ng and about

25 1g of nucleic acid or protein, as applicable. In another embodiment, the effective amount contains between about 10ng and about 100mg of nucleic acid or protein, as applicable. In a further embodiment, the effective amount contains between about 100ng and about

30 10mg of the nucleic acid or protein, as applicable. In a yet a further embodiment, the effective amount contains between about 1 μ g and about 1mg of the nucleic acid or protein, as applicable. In a yet a further embodiment, the effective amount contains between about

10 μ g and about 100 μ g of the nucleic acid or protein, as applicable. In a yet a further embodiment, the effective amount contains between about 100 μ g and about 10mg of the nucleic acid or protein, as applicable.

5

In this invention, administering agents can be effected or performed using any of the various methods and delivery systems known to those skilled in the art. The administering can be performed, for example,
10 intravenously, orally, via implant, transmucosally, transdermally, intramuscularly, and subcutaneously. The following delivery systems, which employ a number of routinely used pharmaceutical carriers, are only representative of the many embodiments envisioned for
15 administering the instant compositions.

Injectable drug delivery systems include solutions, suspensions, gels, microspheres and polymeric injectables, and can comprise excipients such as
20 solubility-altering agents (e.g., ethanol, propylene glycol and sucrose) and polymers (e.g., polycaprylactones and PLGA's). Implantable systems include rods and discs, and can contain excipients such as PLGA and polycaprylactone.

25

Oral delivery systems include tablets and capsules. These can contain excipients such as binders (e.g., hydroxypropylmethylcellulose, polyvinyl pyrrolidone, other cellulosic materials and starch), diluents (e.g.,
30 lactose and other sugars, starch, dicalcium phosphate and cellulosic materials), disintegrating agents (e.g., starch polymers and cellulosic materials) and lubricating agents (e.g., stearates and talc).

Transmucosal delivery systems include patches, tablets, suppositories, pessaries, gels and creams, and can contain excipients such as solubilizers and enhancers (e.g., propylene glycol, bile salts and amino acids),
5 and other vehicles (e.g., polyethylene glycol, fatty acid esters and derivatives, and hydrophilic polymers such as hydroxypropylmethylcellulose and hyaluronic acid).

10 Dermal delivery systems include, for example, aqueous and nonaqueous gels, creams, multiple emulsions, microemulsions, liposomes, ointments, aqueous and nonaqueous solutions, lotions, aerosols, hydrocarbon bases and powders, and can contain excipients such as
15 solubilizers, permeation enhancers (e.g., fatty acids, fatty acid esters, fatty alcohols and amino acids), and hydrophilic polymers (e.g., polycarbophil and polyvinylpyrrolidone). In one embodiment, the pharmaceutically acceptable carrier is a liposome or a
20 transdermal enhancer.

Solutions, suspensions and powders for reconstitutable delivery systems include vehicles such as suspending agents (e.g., gums, zanthans, cellulose and sugars),
25 humectants (e.g., sorbitol), solubilizers (e.g., ethanol, water, PEG and propylene glycol), surfactants (e.g., sodium lauryl sulfate, Spans, Tweens, and cetyl pyridine), preservatives and antioxidants (e.g., parabens, vitamins E and C, and ascorbic acid), anti-
30 caking agents, coating agents, and chelating agents (e.g., EDTA).

In one embodiment of this invention, the delivery system used comprises more than water alone, or more than buffer alone.

5 This invention further provides a method for inhibiting the onset of an inflammatory disorder in a subject comprising administering to the subject a prophylactically effective amount of a nucleic acid which specifically inhibits the expression of RAGE or
10 RAGE G82S in the subject's cells expressing same. This invention further provides a method for treating a subject afflicted with an inflammatory disorder comprising administering to the subject a therapeutically effective amount of a nucleic acid
15 which specifically inhibits the expression of RAGE or RAGE G82S in the subject's cells expressing same.

In the preferred embodiment the subject is human. In a one embodiment the inflammatory disorder is rheumatoid
20 arthritis. In one embodiment the nucleic acid is DNA. In another embodiment the nucleic acid is RNA. In a further embodiment the nucleic acid is an anti-sense nucleic acid that specifically hybridizes to mRNA encoding RAGE or RAGE G82S. In yet a further
25 embodiment the nucleic acid is a catalytic nucleic acid that cleaves mRNA encoding RAGE or RAGE G82S. In yet a further embodiment the nucleic acid is an expressible nucleic acid encoding an anti-sense nucleic acid that specifically hybridizes to mRNA encoding RAGE or RAGE
30 G82S. In yet a further embodiment the nucleic acid is an expressible nucleic acid encoding a catalytic nucleic acid that cleaves mRNA encoding RAGE or RAGE G82S. In one embodiment the subject's cells in which the amount of RAGE or RAGE G82S is to be reduced are

cells which mediate an inflammatory reaction.

This invention further provides a composition comprising a pharmaceutically acceptable carrier and
5 soluble RAGE G82S.

Finally, this invention provides an article of manufacture comprising a packaging material having therein an agent that inhibits binding between RAGE
10 and/or RAGE G82S and a ligand thereof, wherein the packaging material has affixed thereto a label indicating a use for the agent for treating an inflammatory disorder in a subject.

15 In the preferred embodiment of the instant article of manufacture, the subject is human. In one embodiment of the invention, the disorder is rheumatoid arthritis and the agent is selected from the group consisting of soluble RAGE, soluble RAGE G82S, an antibody directed
20 to RAGE, and an antibody directed to RAGE G82S. In one embodiment of the instant article of manufacture, the agent is soluble RAGE. In another embodiment, the agent is soluble RAGE G82S. In a further embodiment, the agent is an antibody directed to RAGE. In yet a
25 further embodiment, the agent is an antibody directed to RAGE G82S, or a small molecule such as an organic molecule.

This invention is illustrated in the Experimental
30 Details section which follows. This section is set forth to aid in an understanding of the invention but is not intended to, and should not be construed to, limit in any way the invention set forth in the claims which follow.

Experimental Details

Murine model of bovine collagen type II-induced arthritis

5 *Induction of arthritis.* Male DBA/1 mice were purchased from the Jackson Laboratories (Bar Harbor, ME). On day 0, mice weighing 20-30 gms were injected intradermally at the base of the tail with bovine type II collagen, 10 200 µg (Sigma), dissolved in acetic acid (0.01M) and emulsified in complete Freund's adjuvant (ICN Biochemcials, Costa Mesa, CA). Three weeks later (day 22), mice were challenged by injection of bovine collagen type II (200 µg) in incomplete Freund's 15 adjuvant (15-17). Beginning three weeks after immunization (day 22), mice were treated with murine soluble RAGE (1) (100 µg per day by intraperitoneal route). Vehicle treatment consisted of murine serum albumin (Sigma), 100 µg per day by intraperitoneal 20 injection, or equal volumes of phosphate-buffered saline (PBS). Treatment was continued daily until sacrifice. All murine studies were performed in accordance with the policies of Columbia University's Institutional Animal Care and Use Committee.

25 *Determination of anti-collagen IgG.* ELISA was performed to determine levels of IgG to bovine type II collagen in dba/1 mice. Plastic Maxisorp 96-well plates (Nunc, Naperville, ILL) were coated with bovine 30 type II collagen (5 µg/ml) in bicarbonate/carbonate buffer; pH 9.3 for 16 hrs at 4°C. Unbound sites were blocked with PBS containing BSA (1%) for 8 hours at 4°C. Wells were washed five times in PBS and then incubated for 16 hrs in samples of mouse plasma 35 (dilution, 1:50) in PBS at 4°C. Wells were washed five

times with PBS and incubated with PBS containing 1% BSA and horse radish peroxidase-labelled anti-mouse IgG (Sigma; 1:500 dilution) for two hours at room temperature. Wells were washed with PBS five times and
5 developed using o-phenylenediamine dihydrochloride (Sigma). A standard curve was prepared using serial dilutions from a mouse antibody cocktail against bovine type II collagen (Chemicon, Temecula, CA).

10 *Assessment of arthritis.* Arthritis was evaluated by clinical and histologic scoring by observers blinded to the experimental conditions. Clinical severity of arthritis in each of four limbs was assessed by the following scale (19): 0=normal; 1=slight erythema and
15 swelling; 2=pronounced edematous swelling; and 3=joint deformity with ankylosis, resulting in a maximum score of 12 per animal. Histologic scoring of arthritis and cartilage and bone destruction was performed on hematoxylin and eosin-stained sections of extremity
20 joints by the following scale (20): (i) synovial lesions: 0, no lesions; 1: mild effect; 2: moderate effect/proliferation; 3: severe lesions with destruction; (ii) cartilage destruction: 0, none; 1, mild; 2, moderate; 3, severe destruction with loss or
25 complete fragmentation of cartilage; and (iii) bone destruction: 0, none; 1, mild destruction of subchondral bone; 2, moderate destruction; and 3, severe destruction with loss of large areas of bone. In other studies, six weeks after initial
30 sensitization, bovine type II collagen (10 µg) was injected into the ear of each mouse. 18 hrs later, thickness of the ear was assessed using calipers by an observer blinded to the experimental conditions.

Immunoblotting and ELISA. Mice were sacrificed 3 or 6 weeks after challenge. The stifle joint was removed and homogenized in tris-buffered saline containing protease inhibitors (Complete Protease Inhibitor, 5 Boehringer-Mannheim, Indianapolis, IN). SDS-PAGE and immunoblotting were performed on extracts of stifle joint tissue using the following antibodies: anti-RAGE IgG and anti-S100/calgranulin IgG as previously described (1) (4.7 and 2.0 µg/ml, respectively). ELISA 10 for TNF-alpha and IL-6 was performed using kits from R&D Systems (Minneapolis, MN).

Zymography. Zymography for detection of MMP-3, 9 and 13 activity was performed using gelatin or casein-laden 15 gels from Novex/Invitrogen (Carlsbad, CA).

Chinese Hamster Ovary (CHO) cell studies

Chinese hamster ovary (CHO) cells were obtained from 20 ATCC (Rockville, MD) and cultured in F12 medium containing fetal bovine serum (10%) (Life Technologies). To generate the 82S allele, the cDNA encoding human RAGE (10) was cloned using the TOPO TA cloning system into pCR2.1TOPO vector for mutagenesis 25 (Invitrogen, Carlsbad, CA). Site-directed mutagenesis to insert the (-82G-) to (-82S-) change was performed using the GeneEditor In Vitro SDM System (Promega, Madison, WI). Sequencing was performed using an ABI310 automated DNA sequencer (Perkin Elmer Biosystems, 30 Foster City, CA) to confirm the inserted sequence changes and to ensure that no other mutations were created. RAGE 82G and 82S cDNAs were excised from pCR2.1TOPO using EcoR I and subcloned into the pcDNA3.1 expression vector (Invitrogen). Cells were transfected

with plasmid DNA using lipofectamine (Life Technologies) encoding the following: pcDNA3.1 containing full-length RAGE cDNA (-82G-), pcDNA3.1 containing RAGE cDNA (-82S-), or pcDNA3.1 containing no insert (mock). 24 hrs after transfection, selection was begun using G418 (1 mg/ml) (Life Technologies). RAGE expression was assessed by immunoblotting in stably-transfected cells after 6 weeks. Cells were incubated with EN-RAGE (Extracellular Newly-Identified RAGE binding protein), a prototypic S100/calgranulin molecule previously-identified to be a signal transduction ligand for RAGE (1), or the indicated mediators. Assessment of activation of phosphorylated MEK 1/2 and p44/p42 MAP kinase by immunoblotting or nuclear translocation of NF-kB was performed. All reagents were tested for endotoxin content using the limulus amebocyte assay (Sigma). Any endotoxin identified in these assays was removed by chromatography of the protein solutions onto Detox-igel columns (Pierce, Arlington Hts, ILL). The material was re-tested in the limulus amebocyte assay for the absence of detectable endotoxin.

Radioligand binding assays. Purified EN-RAGE was radiolabelled using ¹²⁵I and Iodobeads (Pierce, Arlington Heights, ILL) to a specific activity of approximately 5,000 cpm/ng. Radioligand binding assays were performed in 96-well tissue culture dishes containing the indicated transfected CHO cells in the presence of the indicated concentration of radiolabelled EN-RAGE ± an 50-fold molar excess of unlabelled EN-RAGE in PBS containing calcium/magnesium and BSA, 0.2%, for 3 hrs at 37°C. Wells were washed rapidly with washing buffer (PBS containing Tween 20

(0.05%)). Elution of bound material was performed in a solution containing heparin, 1 mg/ml. The aspirate was counted in a gamma counter (LKB, Gaithersburg, MD). Equilibrium binding data were analyzed according to the equation of Klotz and Hunston (57): $B = nKA / 1 + KA$, where B=specifically bound ligand (total binding, wells incubated with tracer alone, minus nonspecific binding, wells incubated with tracer in the presence of excess unlabelled material), n=sites/cell, K=the dissociation constant, and A=free ligand concentration) using nonlinear least-squares analysis (Prism; San Diego, CA). Where indicated, pretreatment with antibodies (70 µg/ml), or human soluble RAGE (50-fold molar excess), was performed.

15 *Activation of pMEK 1/2 and p44/p42 MAP kinases.* CHO cells were incubated with EN-RAGE, 10 µg/ml, for one hr. Cells were lysed in lysis buffer (New England Biolabs, Beverly, MA) and lysate was subjected to centrifugation. Protein concentration of the supernatant was determined by Bio-Rad assay (Bio-Rad, Hercules, CA). Equal amounts of protein were subjected to SDS-PAGE (Novex/Invitrogen). Contents of the gels were transferred to nitrocellulose and immunoblotting performed using anti-phosphorylated/total p44/p42 MAP kinase and MEK 1/2 (New England Biolabs) (1 µg/ml).

Electrophoretic mobility shift assay. Nuclear extracts were prepared using the NE kit (Pierce) and EMSA performed employing consensus ³²P-labelled probe for NF-κB as described (1). Where indicated, cells were treated with either nonimmune/anti-RAGE IgG, or soluble RAGE (1,21).

Peripheral blood-derived mononuclear phagocyte (MPs) studies

Cellular isolation. Venous blood was obtained from
5 healthy volunteers (30 ml) bearing RAGE 82G or 82G/82S
in accordance with the policies of Columbia
University's Institutional Review Board. Due to the
rare numbers of subjects homozygous for the 82S allele,
the studies were confined to those bearing solely 82G
10 or heterozygous 82G/82S. Mononuclear cells were
isolated using Histopaque 1077 (Sigma, St. Louis, MO)
and cultured on plastic dishes for 3 hrs at 37°C.
Nonadherent cells were removed by washing in PBS.
Adherent cells were removed by incubation with EDTA (2
15 mM) for 15 mins at 37°C. Cells (>95% MPs as assessed
by immunostaining with anti-CD68 IgG) were seeded in
tissue culture-coated wells for study.

*Activation of phosphorylated MEK 1/2 and p44/p42 MAP
20 kinases and generation of IL-6 and TNF-alpha.* MPs
were seeded onto the wells of 24-well tissue culture
plates at a density of 5×10^5 cells per well. Cells were
stimulated with either BSA or EN-RAGE (10 µg/ml), and
immunoblotting for detection of phosphorylated/total
25 MEK 1/2 and p44/p42 MAP kinases performed as above.
Supernatant was assayed for IL-6 and TNF-alpha using
ELISA kits from R&D systems (Minneapolis, MN).

Patient population and detection of 82S polymorphism

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RA patients used for association studies meet the
criteria of the American College of Rheumatology (58)
and were taken from patient populations collected
previously at the Medical College of Virginia, Virginia
35 Commonwealth University (31). The following primers

were synthesized for detection of the glycine⁸²serine (G82S) polymorphism of the RAGE gene (8): sense primer: 5' GTAAGCGGGGCTCCTGTTGCA-3' and the antisense primer: 5' GGCCAAGGCTGGGGTTGAAGG-3'. Whole blood (20 µl) was
5 obtained from human volunteers in accordance with the standards and policies of the Institutional Review Boards of the institutions. Genomic DNA was prepared using a kit from Qiagen (Valencia, CA); 10 ng was amplified using Taq DNA polymerase (Life Technologies,
10 Grand Island, NY) in a final volume of 25 µl and PCR performed as follows: 94°C for 30 secs, 62°C for 45 secs, and 72°C for 60 secs for a total of 35 cycles. PCR product (25 µl) was digested with Alu I (Life Technologies), 3U for 16 hr at 37°C, followed by
15 electrophoresis on 2% agarose gels.

Statistical analysis

Statistical comparisons were determined using one-way
20 analysis of variance (ANOVA); where indicated, individual comparisons were performed using students' t-test. For cell culture studies, immunoblotting, zymography and electrophoretic mobility shift assays were performed at least three times. Statistical
25 analyses were performed across each of the experiments by scanning gels/autoradiograms into a densitometer. Arbitrary units of intensity (Image Quant/Molecular Dynamics (Foster City, CA)) were then standardized using untreated cells as control, and assigned a value
30 of "1". Similarly, in animal studies, an arbitrary value of "1" was assigned to bands (immunoblotting or zymography) in assays from animals without collagen-induced arthritis). Statistical analyses to identify the incidence of G82S in human subjects were performed

as previously described (59-60). $P < 0.05$ was considered statistically significant.

Results

5

RAGE and Murine Collagen-Induced Arthritis

As a first test of the hypothesis that RAGE contributed to inflammatory and tissue-destructive mechanisms in arthritis, arthritis was induced in DBA/1 mice by sensitization and challenge with bovine type II collagen, the predominant protein of articular cartilage (15-17). Bovine type II collagen was emulsified in complete Freund's adjuvant and injected intradermally at the base of the tail (day 0). Three weeks later, mice were challenged with bovine type II collagen (day 22). The contribution of RAGE to the pathogenesis of arthritis was studied by treating animals with soluble (s) RAGE (1,18), the extracellular ligand-binding domain of the receptor, 100 μ g per day, beginning three weeks after initial immunization (day 22). In previous studies, blockade of RAGE at this dose effected the greatest decrease in the proinflammatory phenotype in a murine model of delayed-type hypersensitivity (1).

The relevance of RAGE-ligand interaction to murine collagen-induced arthritis was underscored by the increased expression of RAGE and S100/calgranulins in joint tissue. Six weeks after immunization, RAGE and S100/calgranulin expression was increased in joints from mice with arthritis compared with controls, especially within the proliferating synovium (RAGE, fig. 1e&d; S100/calgranulin, fig. 1h&g, respectively).

By immunoblotting, levels of RAGE antigen were enhanced ~2.2-fold in arthritis versus control joints (fig. 1m). Although levels of S100/calgranulins were quite low in joint tissue of control DBA/1 mice without collagen-induced arthritis, expression of these proinflammatory mediators was induced in murine serum albumin-treated mice immunized/challenged with bovine type II collagen (fig. 1n). In mice subjected to blockade of RAGE with sRAGE, levels of RAGE and S100/calgranulin antigens by immunoblotting were significantly reduced compared to mice treated with vehicle (fig. 1m&n, respectively).

In parallel with reduced expression of RAGE and S100/calgranulins in sRAGE-treated joints, clinical indices of inflammation (18) were consistently lower in mice treated with sRAGE vs PBS or murine serum albumin (fig. 2a). In addition, histologic scoring (19) revealed marked decreases in synovial hyperplasia/hypertrophy in the joints of sRAGE-treated mice (fig. 2b). Further, histologic indices of cartilage and bone destruction (20) were significantly decreased in sRAGE-treated mice compared with those animals receiving PBS or murine serum albumin (fig. 2b). To determine if blockade of RAGE suppressed immune/inflammatory responses to bovine type II collagen at extra-articular sites, mice receiving vehicle or sRAGE were injected with bovine type II collagen (10 µg) into ear tissue six weeks after immunization (day 42). Although baseline ear thickness was identical in all groups, 18 hrs after injection of collagen, mice receiving murine serum albumin or PBS revealed significant increases in ear thickness compared with those mice treated with sRAGE (fig. 2c). Levels of serum total IgG developed against bovine type

II collagen were unchanged in the presence of sRAGE vs PBS or murine serum albumin, thereby indicating that blockade of RAGE did not suppress primary immunization (fig. 2d).

5

To dissect the molecular mechanisms underlying the protection afforded by RAGE blockade, levels of cytokines and matrix metalloproteinases (MMPs) in the tissues were assessed. ELISA on extracts prepared from
10 stifle joints revealed a significant reduction in levels of TNF-alpha (~4.3-fold) and IL-6 (~3.1-fold) from mice immunized with bovine type II collagen treated with sRAGE vs murine serum albumin (fig. 3a&b, respectively). Similarly, levels of plasma TNF-alpha
15 and IL-6 antigens were significantly reduced, ~3.5- and 3.6-fold, in sRAGE-treated mice compared with those animals receiving murine serum albumin (fig. 3c&d, respectively).

20 As induction of TNF-alpha and other inflammatory cytokines sets in motion events leading to activation of latent/proenzyme MMPs (21), activity of MMPs 3, 9 and 13 in joint tissue was assessed. Compared with control joints, tissue retrieved from murine serum albumin-
25 treated mice undergoing the collagen-induced arthritis protocol revealed an ~2.5-fold, ~4.0-fold and ~5.3-fold increase in activity of MMPs 3, 9 and 13, respectively, by zymography (fig. 3e&f). That activation of RAGE was critical in this process was demonstrated by the
30 significant reduction in activation of these MMPs in joint tissue of sRAGE-treated mice (fig. 3e&f, respectively).

Taken together, these observations suggested that interaction of S100/calgranulins and RAGE, both enriched in joint tissues in arthritis, activates proinflammatory and tissue-destructive mediators, thereby supporting roles for these molecules in the evolution of joint destruction.

RAGE 82S Allele Enhances Binding/Signalling/Gene Expression by S100/calgranulins

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Polymorphisms within the RAGE gene may influence these effects by altering proinflammatory pathways. To study this, whether the glycine to serine polymorphism (G82S) within the ligand-binding domain of RAGE displayed enhanced ligand affinity and activation of signal transduction pathways was tested. Stably-transfected Chinese hamster ovary (CHO) cells bearing the two RAGE alleles of interest were prepared, either the common RAGE 82G allele, or the variant RAGE 82S allele. CHO cells provide a convenient model, as they are devoid of detectable RAGE prior to (not shown), or after stable transfection with pcDNA3.1 vector alone (mock) (fig. 4a). Stably-transfected CHO cells were made with pcDNA3.1 containing either RAGE 82G or RAGE 82S (fig. 4a). ELISA studies demonstrated that levels of RAGE antigen were identical in both CHO cells transfected to express RAGE 82G or RAGE 82S (data not shown). Radioligand binding studies showed dose-dependent binding of ^{125}I -EN-RAGE (a prototypic S100/calgranulin) to CHO cells expressing RAGE 82G ($\sim 122 \pm 31$ nM) and 82S receptor ($K_d \sim 77 \pm 21$ nM), though the affinity of binding was greater with the 82S allele ($p=0.008$; fig. 4b). In contrast, CHO cells stably transfected with the empty vector displayed no specific binding of ^{125}I -EN-RAGE (fig. 4b). That the interaction of RAGE-bearing CHO

cells with ^{125}I -EN-RAGE was specific for interaction with RAGE was shown by suppression of specific binding in the presence of excess soluble extracellular domain of the receptor (sRAGE), or anti-RAGE IgG, but not by
5 bovine serum albumin (BSA) or nonimmune IgG (fig. 4c).

These observations led to testing the concept that the 82S RAGE isoform might amplify cellular activation beyond that seen in cells bearing wild-type RAGE.
10 Incubation of mock-transfected CHO cells with a physiologically-relevant concentration of EN-RAGE (4), 10 $\mu\text{g/ml}$, did not significantly increase intensity of the bands corresponding to phosphorylated MEK 1/2 or p44/42 MAP kinases (22-23) (figs. 4d&e, respectively;
15 lanes 1-5). However, exposure of RAGE 82G CHO transfectants to EN-RAGE increased by ~ 2.2 -fold and ~ 1.9 -fold phosphorylated MEK 1/2 and p44/42 MAP kinases, compared with cultures incubated with BSA alone ($p < 0.01$; fig. 4d&e, respectively; lanes 7&6).
20 CHO transfectants bearing RAGE 82S incubated with EN-RAGE displayed ~ 3.6 -fold and ~ 4.1 -fold increase in phosphorylated MEK 1/2 and p44/42 compared with BSA ($p < 0.01$; fig. 4d&e, respectively; lanes 12&11). Thus, compared with cells expressing the common RAGE 82G
25 allele, CHO cells expressing the variant RAGE 82S displayed significantly increased phosphorylation of MEK 1/2 and p44/p42 MAP kinases by ~ 1.6 - and ~ 2.2 -fold ($p < 0.05$; fig. 4d&e, respectively; lanes 12&7). In both RAGE 82G and RAGE 82S-transfected cells, cellular
30 activation by EN-RAGE was due to engagement of RAGE as demonstrated by suppression of phosphorylation of MEK 1/2 and p44/p42 by excess sRAGE (fig. 4d&e, respectively; lanes 8&13), or anti-RAGE IgG (fig. 4d&e, respectively, lanes 9&14). Nonimmune IgG was without

effect (fig. 4d&e, respectively; lanes 10&15). In each case, respective levels of total MEK 1/2 and p44/p42 MAP kinases were identical (data not shown).

5 To further support the hypothesis that RAGE 82S enhanced activation of key proinflammatory signal transduction pathways, nuclear translocation of NF- κ B in CHO transfectants exposed to EN-RAGE were assessed. Electrophoretic mobility shift assays (EMSA) using ³²P-
10 labelled consensus NF- κ B probe and nuclear extracts from mock-transfected CHO cells showed no increase in intensity of the gel shift band after cultures were exposed to EN-RAGE (fig. 4f, lane 2). In CHO transfectants expressing RAGE 82G, there was a
15 prominent ~2.4-fold increase in intensity in nuclear binding activity following incubation of cultures with EN-RAGE compared to BSA ($p < 0.05$; fig. 4f, lanes 7&6, respectively). NF- κ B activation was even more striking when RAGE 82S was substituted for RAGE 82G. RAGE 82S
20 CHO transfectants displayed ~5.1-fold increased intensity of the gel shift band consequent to the presence of EN-RAGE, compared to incubation with BSA ($p < 0.001$; fig. 4f, lanes 12&11, respectively). Thus, RAGE-mediated NF- κ B activation due to EN-RAGE was
25 significantly enhanced by ~2.1-fold comparing RAGE 82S to 82G; $p < 0.01$. That activation of NF- κ B in transfected CHO cells by EN-RAGE resulted from ligation of common or variant RAGE was confirmed by its suppression in the presence of sRAGE (fig. 4f, lanes
30 8&13, respectively), or anti-RAGE IgG (fig. 4f, lanes 9&14, respectively), but not by nonimmune IgG (fig. 4f, lanes 10&15, respectively).

A critical test of the concept that S100/calgranulin-RAGE 82S interaction might enhance proinflammatory effects was whether mononuclear phagocytes (MPs) retrieved from human subjects bearing the variant RAGE 82S allele displayed heightened activation of signal transduction molecules in the presence of ligand. Due to the rare number of subjects homozygous for 82S in our study populations, studies were limited to those individuals homozygous for the common RAGE allele 82G, or those subjects heterozygous for 82G and 82S alleles. ELISA revealed that basal levels of RAGE antigen did not differ between MPs bearing 82G or 82G/82S (data not shown). Signalling was compared in MPs from subjects bearing RAGE 82G and RAGE 82G/82S by assessing activation of MEK 1/2 and p44/p42 MAP kinases. In the presence of EN-RAGE, MPs isolated from individuals bearing RAGE 82G/82S displayed an ~4.5- and 4.3-fold increase in phosphorylated MEK 1/2 and p44 and p42 MAP kinases, respectively, compared with unstimulated cells ($p < 0.01$; fig. 5a,b,c&d). However, MPs bearing the common RAGE 82G allele exposed to EN-RAGE revealed a significant, although smaller (~2.6- and 2.3-fold) increase in activation of MEK 1/2 and p44/p42 MAP kinases, respectively ($p < 0.01$; fig. 5a,b,c&d). Importantly, EN-RAGE-triggered activation of MEK 1/2 and p44/p42 MAP kinases was significantly increased in heterozygous RAGE 82G/82S-bearing MPs vs those MPs expressing RAGE 82G ($p < 0.05$; fig. 5b&d).

To assess the functional consequences of enhanced binding and activation of signal transduction molecules stimulated upon ligation of RAGE by EN-RAGE, production of key inflammatory and tissue-degradative mediators linked to RA (24-26) by MPs bearing common or variant

RAGE alleles were examined. Exposure of RAGE 82G-bearing MPs to EN-RAGE caused a significant increase in generation of TNF-alpha detected in culture supernatant compared with quiescent cultures (470 ± 53 vs 15 ± 2 pg/ml; $p < 0.001$) (fig. 5e). However, upon incubation of human MPs bearing RAGE 82G/82S, an even greater increase in elaborated TNF-alpha was observed in culture supernatants compared to basal levels (850 ± 64 vs 17 ± 2 pg/ml; $p < 0.01$) (fig. 5e). Importantly, although basal levels of TNF-alpha did not differ between RAGE 82G and 82G/82S-bearing MPs, levels of TNF-alpha in EN-RAGE-stimulated MPs were increased ~1.8-fold in the presence of RAGE 82G/82S compared with RAGE 82G ($p < 0.01$; fig. 5e). Similarly, RAGE 82G-bearing MPs exposed to EN-RAGE displayed a small, but significant increase in generation of IL-6 compared with basal expression (55 ± 7 vs 15 ± 2 pg/ml; $p < 0.01$) (fig. 5f). However, MPs bearing RAGE 82G/82S revealed augmented generation of IL-6 upon incubation with EN-RAGE compared with unstimulated controls (160 ± 14 vs 12 ± 2 pg/ml; $p < 0.01$) (fig. 5f). EN-RAGE-stimulated MPs bearing RAGE 82G/82S generated increased amounts of IL-6 compared with cells from subjects bearing RAGE 82G (~2.9-fold) ($p < 0.01$; fig. 5f).

25

A central means by which structural elements of joints are degraded in unchecked RA is by generation of MMPs, such as MMP-9 (27-30). It was hypothesized that RAGE-mediated MP activation would augment generation of MMP activity on cells bearing RAGE 82G/82S versus RAGE 82G. Activity of MMP-9 in MPs retrieved from subjects bearing the common or variant RAGE alleles was tested. MPs from subjects bearing RAGE 82G displayed an ~2.3-fold increase in MMP-9 activity in the presence of EN-

RAGE compared with basal expression ($p < 0.01$; fig. 5g&h). However, MPs isolated from individuals bearing RAGE 82G/82S demonstrated an ~4.3-fold increase in EN-RAGE-mediated MMP-9 activity compared with baseline (5 $p < 0.01$; fig. 5g&h). Although basal levels of MMP-9 activity did not differ among 82G- or 82G/82S-bearing MPs, the extent of EN-RAGE-mediated enhanced MMP-9 activity was significantly enhanced, ~1.9-fold, in MPs bearing RAGE 82G/82S vs RAGE 82G ($p < 0.05$; fig. 5g&h).

10

Taken together, the studies in DBA/1 mice suggest that activation of RAGE amplifies cytokine generation and activation of MMPs in joint tissue primed by immunization with bovine type II collagen. These (15 considerations, together with the findings in vitro suggesting that S100/calgranulin-RAGE interaction modifies cellular responses, especially in the presence of the 82S allele, led to the evaluation of the association of the RAGE 82S allele with RA.

20

RAGE 82S Allele and Association with Rheumatoid Arthritis

Subjects from a previously-reported Caucasian (25 population of RA patients ($n=205$) and matched controls ($n=169$) (31) were evaluated. The RAGE 82S allele exhibited an association with RA with relative risk (RR) =2.6, $p < 0.001$ (table 1a). Further analyses were performed to account for the linkage disequilibrium (30 described previously between RAGE 82S and an RA-associated allele, HLA-DRB*0401 (9,32-34). To assess whether the RAGE 82S allele confers risk independent of DRB1*0401, the frequency of the RAGE 82S allele in 0401+ patients and controls was compared. This analysis (35 did not reveal a significant independent effect of RAGE

82S on RA disease susceptibility; $p=0.13$ by Fisher's exact test (table 1b). However, DRB1*0401 continued to exhibit a significant association with RA in the absence of the RAGE 82S allele (table 1c).

5

Discussion

S100/calgranulin molecules have been extensively linked to immune/inflammatory diseases. Although their
10 intracellular functions are relatively well-defined, such as calcium-dependent chemotaxis, degranulation and phagocytosis by activated granulocytes (5-6,35), an increasingly recognized view is that these molecules may exhibit distinct functions in the extracellular
15 milieu (35). These concepts are consistent with the observations that the degree of elevation of S100/calgranulins in inflamed foci may mirror the extent of disease activity. For example, in addition to their association with disease severity in RA (4),
20 S100/calgranulins have been linked to disease progression in inflammatory bowel disease (36,37). Furthermore, strongly increased expression of a member of this family of molecules, psoriasin, has been demonstrated in psoriatic lesions compared with
25 adjacent unaffected skin in human subjects (38), thus further supporting the premise that S100/calgranulins may not simply be "innocent bystanders," but, rather, by virtue of their engagement of RAGE (1), active participants in the host inflammatory response. One
30 can speculate that triggered by interaction with accumulating S100/calgranulins, activation of RAGE engages key signalling pathways linked to proinflammatory responses, such as MAP kinases and NF- κ B, thereby amplifying expression of molecules linked

to chronic cellular perturbation and tissue destruction. This hypothesis appears to be supported by the studies in DBA/1 mice immunized/challenged with bovine type II collagen, as activation of RAGE mediated, at least in part, increased generation of inflammatory cytokines and activation of MMPs. Indeed, blockade of the receptor was associated with decreased clinical and histologic indices of inflammation and bone/cartilage destruction in this model.

10

Polymorphic differences in the RAGE gene may lead to varied effects of RAGE in inflammatory settings. This led to the investigation of the effects of the G82S polymorphism in the binding domain of RAGE on receptor function and on the inflammatory response. In transfected CHO cells, increased ligand affinity was found to be conferred by the 82S isoform, which, upon further analysis, demonstrated upregulation of intracellular signalling pathways linked to modulation of proinflammatory genes upon interaction with S100/calgranulin. Using isolated MPs from human subjects, significant differences in inflammatory responses between cells isolated from homozygous wild-type G82G and heterozygous G82S subjects were demonstrated. These effects were quite marked between wild-type and heterozygote subjects, thereby suggesting that the 82S allele alone may impart an even greater effect.

30 Finally, the prevalence of the 82S allele in a population of subjects with RA was investigated. The results demonstrated a statistically-significant increase in prevalence of the 82S allele in subjects with RA. Sub-analysis for the presence/absence of

DRB1*0401 led to the loss of an statistically-significant association of 82S with RA ($p=0.13$ despite odds ratio 3.2). Conversely, DRB1*0401 continued to exhibit a significant association with RA in the
5 absence of the RAGE 82S allele. Clearly, the large genetic susceptibility to RA related to the MHC (39-40) cannot be ascribed solely to the RAGE 82S allele. Due to the relatively low frequency of the 82S allele and low subject numbers in these Caucasian cases and
10 controls after sub-analysis, it remains possible that the RAGE 82S polymorphism is related independently to RA.

The literature is conflicting with respect to a role
15 for DR4 haplotypes in disease severity (41-50). In view of the linkage disequilibrium between DRB1*0401 and the RAGE 82S allele, it will be interesting to specifically test whether DR4 haplotypes containing the RAGE 82S allele have an influence on disease progression and
20 outcome. By analogy with the observations in cultured CHO and human mononuclear cells, one can speculate it may be possible to identify a subset of subjects with RA particularly vulnerable to the sequelae of heightened production/activation of proinflammatory
25 mediators. Specifically, in parallel with outcome measures of clinical severity, such as numbers of bony erosions, the extent of elevation of TNF-alpha, IL-6 and MMP activities in rheumatoid joint fluid and/or RA plasma must be correlated with subject genotype.

30

A general implication of these studies is that the classical DRB1 associations with RA may not completely explain the underlying reasons for the strong linkage and association of the MHC with RA. Indeed, a number

of recent reports indicate that additional genes in the TNF region may influence disease susceptibility, independent of DRB1, and unrelated to DR4 haplotypes (51-52). A full understanding of the role of the MHC in RA and other autoimmune diseases will require a more detailed analysis of this genetic region, since there are many genes within the MHC, in addition to RAGE, which may influence the immune/inflammatory response (53).

10

Lastly, in addition to the 82S variant of the RAGE gene, it will be important to examine the potential role of other variants of RAGE in disease susceptibility and/or severity in immune/inflammatory diseases. In this context, recently identified polymorphisms within the promoter of the gene encoding RAGE may hold the key to unique elements controlling enhanced transcription of the RAGE gene and increased expression of cell surface RAGE (54). As a number of studies have suggested that expression of RAGE is increased at sites of ligand accumulation such as diabetic atherosclerotic lesions, infected periodontium or Alzheimer disease brain (13,18,55), polymorphisms within regulation elements and/or ligand-binding regions may, alone or in combination, orchestrate RAGE's availability and/or functional potential in a given milieu (56).

In conclusion, one can propose that these observations in an animal model of inflammatory arthritis and in S100/calgranulin-stimulated activation of 82S RAGE-bearing cells highlight ligand-RAGE engagement as an amplifier of proinflammatory mechanisms in immune/inflammatory diseases. These findings provide a

compelling rationale to expand the analysis of RAGE polymorphisms and investigate their potential contribution to susceptibility and/or progression of the inflammatory response in diverse diseases with an
5 immune/inflammatory component, such as RA, atherosclerosis and diabetes.

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What is claimed is:

1. A method for inhibiting binding between RAGE and a ligand thereof comprising contacting the RAGE with an agent selected from the group consisting of soluble RAGE, soluble RAGE G82S, and an antibody directed to RAGE.
2. The method of claim 1, wherein the agent is soluble RAGE.
3. The method of claim 1, wherein the agent is soluble RAGE G82S.
4. The method of claim 1, wherein the agent is an antibody directed to RAGE.
5. A method for inhibiting binding between RAGE G82S and a ligand thereof comprising contacting the RAGE G82S with an agent selected from the group consisting of soluble RAGE, soluble RAGE G82S, and an antibody directed to RAGE G82S.
6. The method of claim 5, wherein the agent is soluble RAGE.
7. The method of claim 5, wherein the agent is soluble RAGE G82S.
8. The method of claim 5, wherein the agent is an antibody directed to RAGE G82S.
9. A method for treating an inflammatory disorder in a subject comprising administering to the subject

a therapeutically effective amount of an agent that inhibits binding between RAGE and/or RAGE G82S and a ligand thereof.

- 5 10. The method of claim 9, wherein the inflammatory disorder is rheumatoid arthritis.
11. The method of claim 9, wherein the subject is a human.
- 10 12. The method of claim 9, wherein the agent is selected from the group consisting of soluble RAGE, soluble RAGE G82S, an antibody directed to RAGE, and an antibody directed to RAGE G82S
- 15 13. The method of claim 12, wherein the agent is soluble RAGE.
14. The method of claim 12, wherein the agent is soluble RAGE G82S.
- 20 15. The method of claim 12, wherein the agent is an antibody directed to RAGE.
- 25 16. The method of claim 12, wherein the agent is an antibody directed to RAGE G82S.
17. A method for inhibiting the onset of an inflammatory disorder in a subject comprising administering to the subject a prophylactically effective amount of an agent that inhibits binding between RAGE and/or RAGE G82S and a ligand thereof.
- 30

18. The method of claim 17, wherein the subject is a human.
19. The method of claim 17, wherein the agent is
5 selected from the group consisting of soluble RAGE, soluble RAGE G82S, an antibody directed to RAGE, and an antibody directed to RAGE G82S.
20. The method of claim 19, wherein the agent is
10 soluble RAGE.
21. The method of claim 19, wherein the agent is soluble RAGE G82S.
22. The method of claim 19, wherein the agent is an
15 antibody directed to RAGE.
23. The method of claim 19, wherein the agent is an
20 antibody directed to RAGE G82S.
24. A method for inhibiting the onset of an inflammatory disorder in a subject comprising administering to the subject a prophylactically effective amount of a nucleic acid which
25 specifically inhibits the expression of RAGE or RAGE G82S in the subject's cells expressing same.
25. A method for treating a subject afflicted with an inflammatory disorder comprising administering to
30 the subject a therapeutically effective amount of a nucleic acid which specifically inhibits the expression of RAGE or RAGE G82S in the subject's cells expressing same.

26. The method of claim 24 or 25, wherein the nucleic acid is DNA.
27. The method of claim 24 or 25, wherein the nucleic acid is RNA.
28. The method of claim 24 or 25, wherein the nucleic acid is an anti-sense nucleic acid that specifically hybridizes to mRNA encoding RAGE or RAGE G82S.
29. The method of claim 24 or 25, wherein the nucleic acid is a catalytic nucleic acid that cleaves mRNA encoding RAGE or RAGE G82S.
30. The method of claim 24 or 25, wherein the nucleic acid is an expressible nucleic acid encoding an anti-sense nucleic acid that specifically hybridizes to mRNA encoding RAGE or RAGE G82S.
31. The method of claim 24 or 25, wherein the nucleic acid is an expressible nucleic acid encoding a catalytic nucleic acid that cleaves mRNA encoding RAGE or RAGE G82S.
32. The method of claim 24 or 25, wherein the subject's cells in which the amount of RAGE or RAGE G82S is to be reduced are cells which mediate an inflammatory reaction.
33. The method of claim 24 or 25, wherein the subject is human.

34. The method of claim 24 or 25, wherein the inflammatory disorder is rheumatoid arthritis.
- 5 35. A composition comprising a pharmaceutically acceptable carrier and soluble RAGE G82S.
- 10 36. An article of manufacture comprising a packaging material having therein an agent that inhibits binding between RAGE and/or RAGE G82S and a ligand thereof, wherein the packaging material has affixed thereto a label indicating a use for the agent for treating an inflammatory disorder in a subject.
- 15 37. The article of claim 36, wherein the subject is a human.
- 20 38. The article of claim 36, wherein the inflammatory disorder is rheumatoid arthritis and the agent is selected from the group consisting of soluble RAGE, soluble RAGE G82S, an antibody directed to RAGE, and an antibody directed to RAGE G82S.
- 25 39. The article of claim 38, wherein the agent is soluble RAGE.
40. The article of claim 38, wherein the agent is soluble RAGE G82S.
- 30 41. The article of claim 38, wherein the agent is an antibody directed to soluble RAGE.
42. The article of claim 38, wherein the agent is an antibody directed to soluble RAGE G82S.

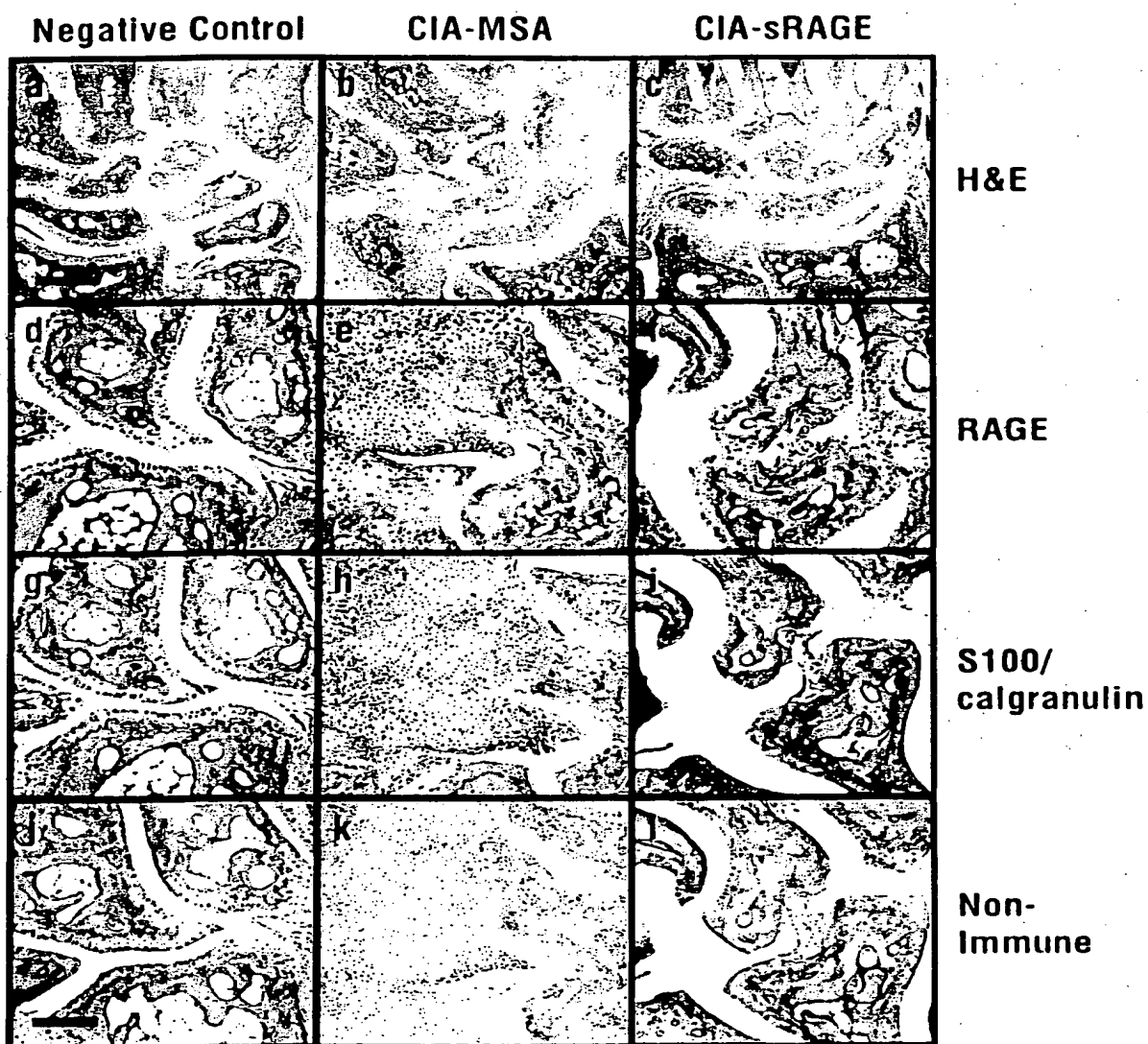
RAGE AND RAGE G82S-RELATED METHODS AND COMPOSITIONS FOR
TREATING INFLAMMATORY DISORDERS

Abstract of the Disclosure

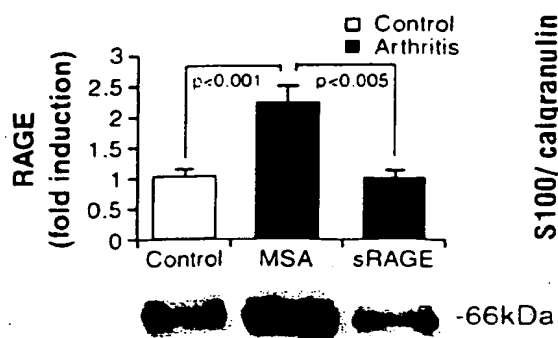
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This invention provides methods, compositions and
articles of manufacture for inhibiting the onset of and
treating inflammatory disorders such as rheumatoid
arthritis. The instant invention is based on the
10 blockade of RAGE and/or RAGE G82S function.

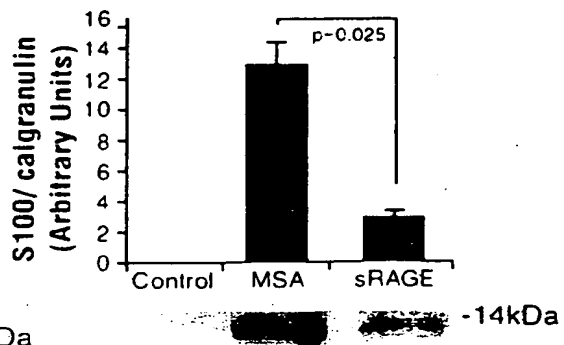
Figures 1A-1N



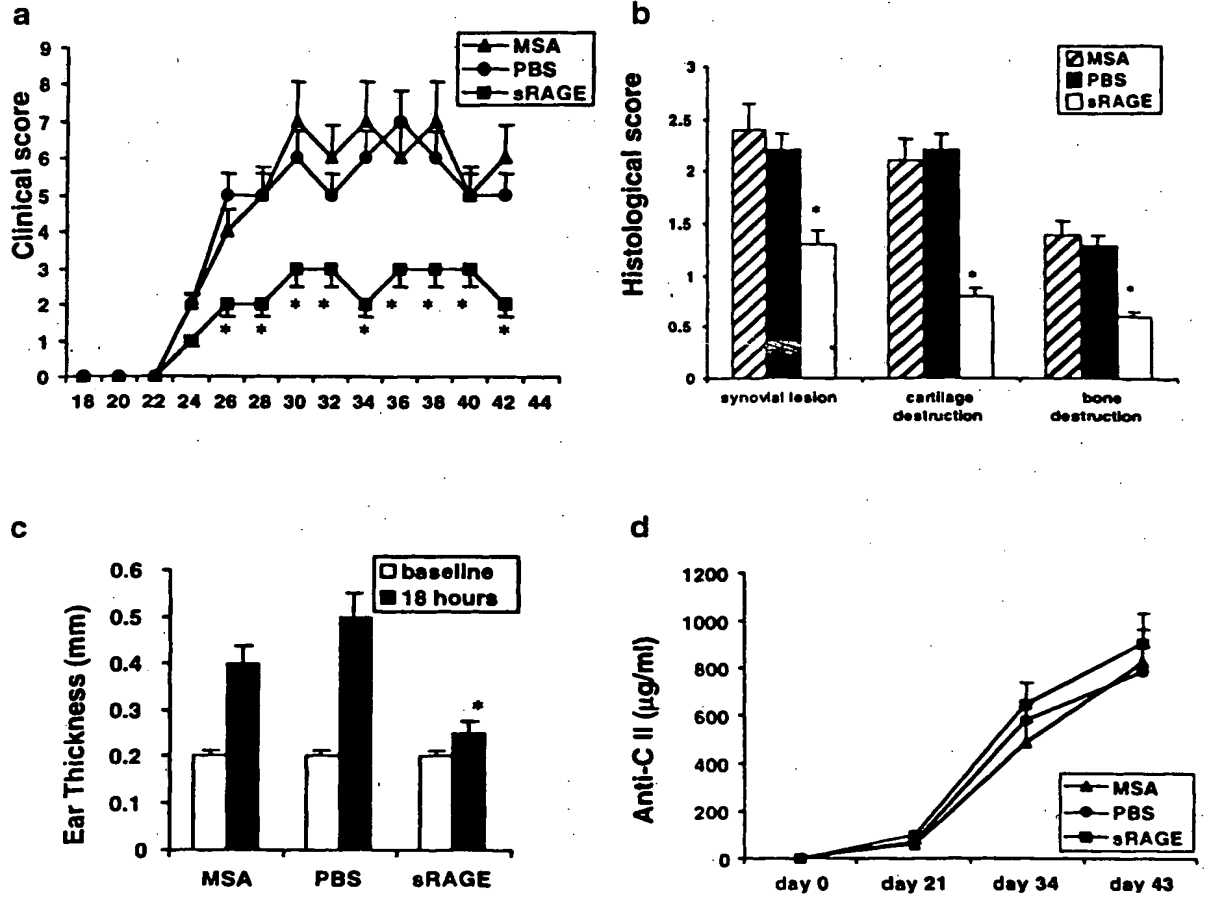
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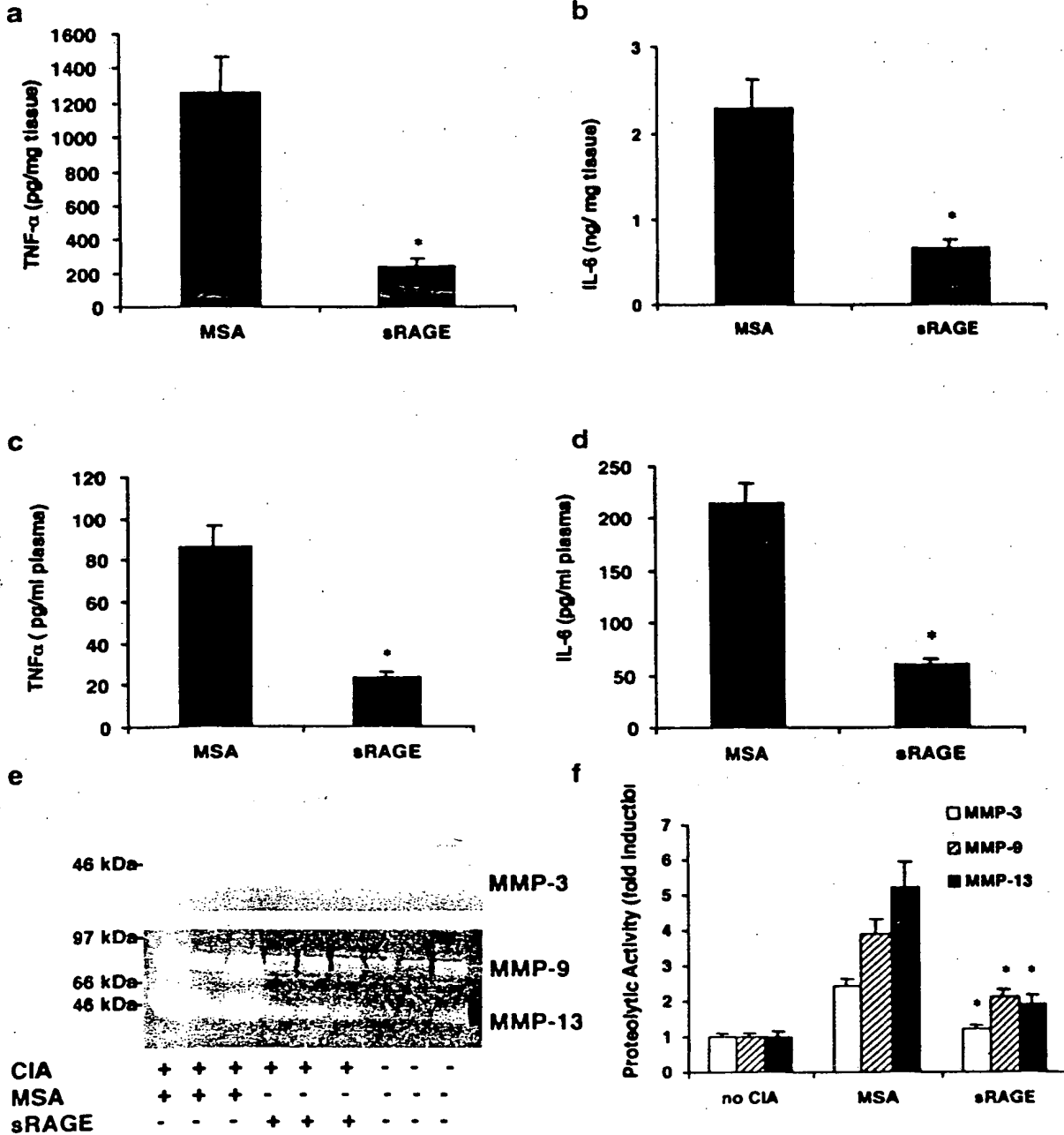
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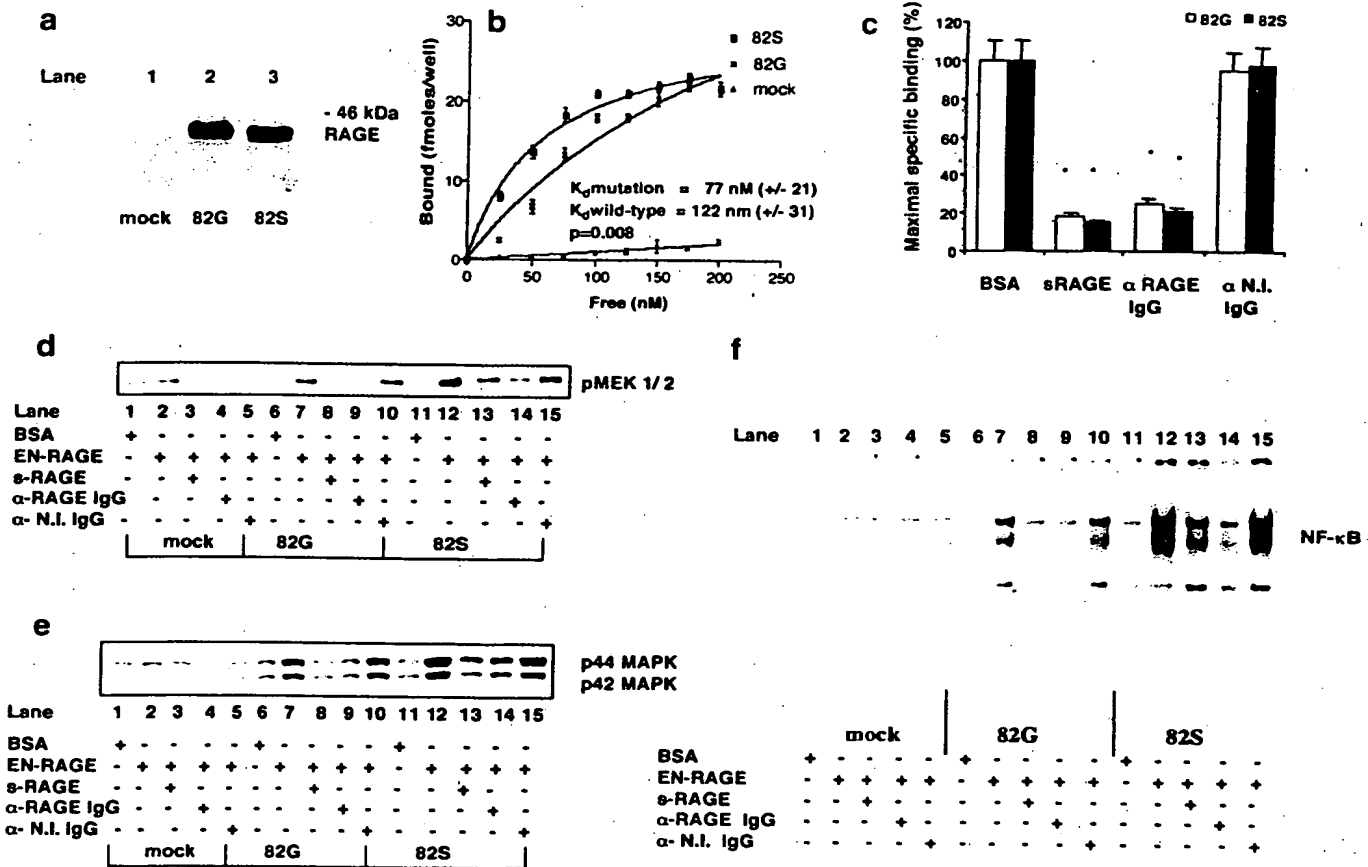
Figures 2A-2D



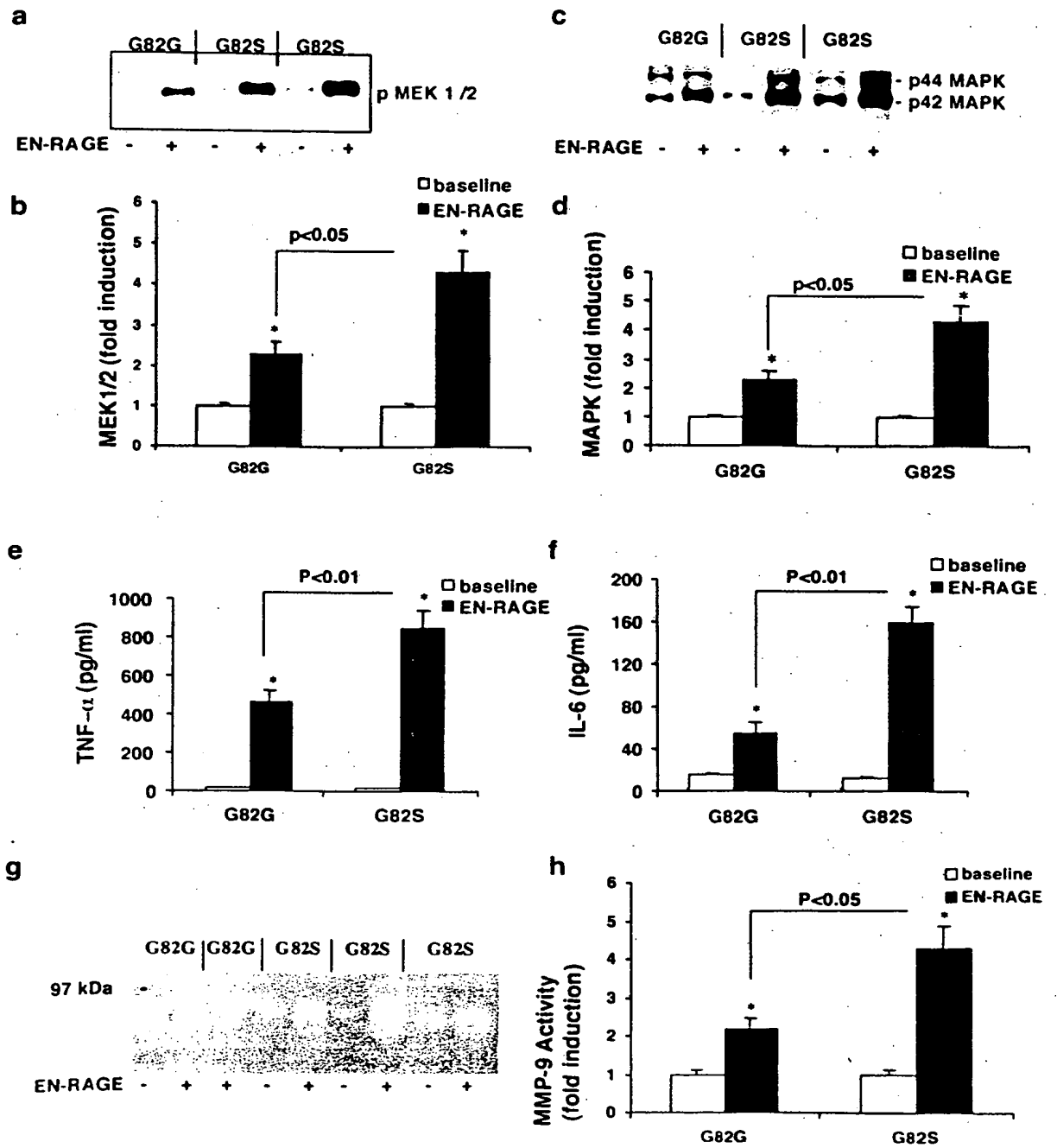
Figures 3A-3F



Figures 4A-F



Figures 5A-5H



*Application
for
United States Letters Patent*

To all whom it may concern:

Be it known that

Ann Marie Schmidt and David M. Stern

have invented certain new and useful improvements in

**RAGE G82S-RELATED METHODS
AND COMPOSITIONS FOR TREATING
INFLAMMATORY DISORDERS**

of which the following is a full, clear and exact description.

**RAGE G82S-RELATED METHODS AND COMPOSITIONS FOR TREATING
INFLAMMATORY DISORDERS**

5

This application claims the benefit of U.S. Provisional Application No. 60/469,428, filed May 9, 2003, the contents of which are hereby incorporated by reference.

10 Throughout this application, various publications are referenced. Full citations for these publications may be found immediately preceding the claims. The disclosures of these publications are hereby incorporated by reference into this application in
15 order to more fully describe the state of the art as of the date of the invention described and claimed herein.

Background of the Invention

20 The Receptor for Advanced Glycation Endproducts ("RAGE"), a multi-ligand member of the immunoglobulin superfamily of cell surface molecules, has been implicated in amplification of proinflammatory responses (1). S100/calgranulins, proinflammatory
25 signal transduction ligands of the receptor, are enriched in joints of subjects with rheumatoid arthritis ("RA") (2-4). Members of this family of molecules, long-associated with classic immune/inflammatory disorders (5-6), may be released by
30 activated inflammatory effector cells such as monocytes (7), thereby freeing them to engage cell surface RAGE and amplify host inflammatory responses. Indeed, accumulation of S100/calgranulins in synovial fluid and plasma of subjects with RA has been correlated with
35 indices of disease severity, such as bony erosions (4).

Recent studies have highlighted the possibility that polymorphisms within key domains of RAGE may influence its function, so that under conditions of increased
5 ligand accumulation, individuals may be predisposed to heightened inflammatory responses (8-9). Importantly, a polymorphism of the RAGE gene has been identified within the V-type immunoglobulin domain in the extracellular region of the receptor, consisting of a
10 glycine to serine change at position 82 (8). In previous studies, it was found that the ligands of RAGE, including S100/calgranulins, engage the V-domain of the receptor and activate signal transduction pathways, thereby modulating gene expression (1,10-14).

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Summary of the Invention

This invention provides a method for inhibiting binding between RAGE and a ligand thereof comprising contacting
5 the RAGE with an agent comprising soluble RAGE G82S, a ligand-binding portion of soluble RAGE G82S, or an antibody directed to RAGE G82S.

This invention further provides a method for inhibiting
10 binding between RAGE G82S and a ligand thereof comprising contacting the RAGE G82S with an agent comprising soluble RAGE G82S, a ligand-binding portion of soluble RAGE G82S, or an antibody directed to RAGE G82S.

15 This invention further provides a method for treating an inflammatory disorder in a subject comprising administering to the subject a therapeutically effective amount of an agent that inhibits binding
20 between RAGE G82S and a ligand thereof.

This invention further provides a method for inhibiting the onset of an inflammatory disorder in a subject comprising administering to the subject a
25 prophylactically effective amount of an agent that inhibits binding between RAGE G82S and a ligand thereof.

This invention further provides soluble RAGE G82S or a
30 ligand-binding portion thereof.

This invention further provides a composition comprising a pharmaceutically acceptable carrier and (i) soluble RAGE G82S, (ii) a ligand-binding portion of

soluble RAGE G82S, and/or (iii) a nucleic acid which specifically inhibits the expression of RAGE G82S in a cell expressing same.

5 This invention further provides a method for inhibiting the onset of an inflammatory disorder in a subject comprising administering to the subject a prophylactically effective amount of a nucleic acid which specifically inhibits the expression of RAGE G82S
10 in the subject's cells expressing same.

This invention further provides a method for treating a subject afflicted with an inflammatory disorder comprising administering to the subject a
15 therapeutically effective amount of a nucleic acid which specifically inhibits the expression of RAGE G82S in the subject's cells expressing same.

Finally, this invention provides an article of
20 manufacture comprising a packaging material having therein (i) an agent that inhibits binding between RAGE G82S and a ligand thereof and/or (ii) a nucleic acid which specifically inhibits the expression of RAGE G82S in a cell expressing same, wherein the packaging
25 material has affixed thereto a label indicating a use for the agent and/or nucleic acid for treating an inflammatory disorder in a subject.

30

Brief Description of the Figures

Figures 1A-1N. Induction of arthritis by bovine type II collagen in DBA/1 mice: increased expression of RAGE and S100/calgranulins. Six weeks after immunization (day 42), joint tissue from the hind feet (a-l) or stifle joint (m-n) was prepared for study.

1A-1L. Histology. In a-b-c, tissue was subjected to H&E analysis. Immunohistochemistry using anti-RAGE IgG (d-e-f); anti-S100/calgranulin IgG (g-h-i); or rabbit nonimmune IgG (j-k-l) was performed. Scale bar: a-b-c, 300 μ m; and d-l, 60 μ m.

1M-1N. Immunoblotting. Lysates were prepared from stifle joints and subjected to immunoblotting using anti-RAGE IgG (m); or anti-S100/calgranulin IgG (n). N=5 mice per condition were employed. Densitometric analysis of band intensity from all mice in each group was performed, and the mean \pm SD is shown. Representative bands are shown. Statistical considerations are indicated in the figure.

Figures 2A-2D. Blockade of RAGE suppresses inflammation in DBA/1 mice immunized/challenged with bovine type II collagen.

2A. Clinical score. At the indicated times after immunization with bovine type II collagen and treatment with murine sRAGE or vehicle (murine serum albumin (MSA) or PBS), clinical scoring was performed by blinded observers. The mean \pm standard deviation (SD) is shown; n= 10 mice per group. * indicates $p < 0.05$.

2B. Histologic score. Histologic scoring of joints in four limbs was performed on H&E-stained sections from mice six weeks after immunization. The mean \pm SD is shown; n=10 mice per group. * indicates $p < 0.05$.

5

2C. Ear swelling. Six weeks after immunization with bovine type II collagen, murine serum albumin (MSA)-, PBS- and sRAGE-treated mice were injected with bovine type II collagen into ear tissue. Ear thickness was
10 measured by a blinded observer immediately prior to local injection, and 18 hrs later. The mean \pm SD is shown; n=5 mice per group. * indicates $p < 0.05$ vs ear thickness in PBS- and MSA-treated groups.

15 2D. Antibody response to bovine type II collagen. At the indicated time points after immunization, plasma was retrieved from DBA/1 mice and subjected to ELISA for determination of total serum IgG levels to type II collagen. There were no statistically-significant
20 differences in total IgG levels among the groups.

Figures 3A-3F. Blockade of RAGE suppresses generation of cytokines and MMPs in DBA/1 mice immunized/challenged with bovine type II collagen.

25

3A-3B. Joint tissue. Stifle joint tissue was retrieved six weeks after immunization with bovine type II collagen. Lysates were subjected to ELISA for detection of murine TNF-alpha (a) and IL-6 (b).
30 Results were normalized per mg/tissue extract.

3C-3D. Plasma. Plasma from mice was subjected to ELISA for levels of TNF-alpha (c) and IL-6 (d). In

a-b-c-d, n=10 mice per group. The mean \pm SD is shown.
In a-d, * indicates $p < 0.05$.

3E-3F. Zymography. Stifle joint tissue was retrieved
5 and lysates were prepared and zymography for detection
of activity of MMP-2, -9 or -13 was performed. N=10
mice/group. Densitometric analysis of band intensity
from all mice/groups was performed, and the mean \pm SD
is shown. Representative bands are shown. In f, *
10 indicates $p < 0.05$ vs MSA-treated mice with collagen-
induced arthritis.

Figures 4A-4F. CHO cells bearing RAGE 82S display
increased affinity and cellular responsiveness to EN-
15 RAGE. CHO cells, which endogenously do not express
RAGE antigen, were stably-transfected with pcDNA3.1
vector containing cDNA encoding human RAGE 82G or the
variant allele 82S.

20 4A. Immunoblotting. Lysates of stably-transfected CHO
cells were prepared and subjected to immunoblotting
using anti-human RAGE IgG.

4B-4C. Radioligand binding assays. Purified EN-RAGE
25 was radiolabelled using 125 I and radioligand binding
assays were performed in 96-well tissue culture dishes
containing the indicated transfected CHO cells.
Equilibrium binding data were analyzed according to the
equation of Klotz and Hunston. Where indicated,
30 pretreatment with either antibodies, human soluble RAGE
or bovine serum albumin was performed. The mean \pm
standard deviation (SD) is shown. In c, * indicates
 $p < 0.01$ versus respective controls.

4D-4E. Activation of MEK 1/2 and p44/p42 MAP kinases. The indicated stably-transfected CHO cells were incubated with EN-RAGE for one hr. Immunoblotting of cell lysates was performed using anti-phosphorylated
5 MEK 1/2 (d) or p44/p42 MAP kinase (e). Where indicated, pretreatment with either BSA or sRAGE, or the indicated IgG for 2 hrs was performed. Control immunoblotting using antibody to total MEK 1/2 and p44/p42 MAP kinase revealed that there were no
10 differences in levels of total MEK 1/2 or p44/p42 (not shown).

4F. Activation of NF- κ B. Nuclear extracts were prepared from the indicated stably-transfected CHO
15 cells incubated with EN-RAGE for 6 hrs and EMSA was performed. Where indicated, cells were pretreated with either nonimmune/anti-RAGE IgG, soluble RAGE or BSA for 2 hrs prior to incubation with EN-RAGE. In d-e-f, bands were scanned into a densitometer, and band
20 density was quantified using ImageQuant. These experiments were performed at least three times and subjected to statistical analyses as indicated in the text and the figure legend. Representative experiments are shown.

25

Figures 5A-5H. Human peripheral blood mononuclear phagocytes (MPs) expressing RAGE 82S display increased responsiveness to EN-RAGE. MPs were purified from the blood of human subjects bearing RAGE (G82G) (n=5) or
30 heterozygotes (G82S) (n=5).

5A-5D. Activation of MEK 1/2 and p44/p42 MAP kinases. The indicated MPs were incubated with EN-RAGE or no mediator for one hr. Cell lysates were prepared and

immunoblotting performed using anti-phosphorylated MEK 1/2 (a,b) or p44/p42 MAP kinase (c,d). Densitometric analysis was performed and mean \pm SD is shown in b and d. Immunoblotting using antibody to total MEK 1/2 and p44/p42 MAP kinase revealed that there were no differences in levels of total MEK 1/2 or p44/p42 (not shown).

5E-5F. Generation of TNF-alpha and IL-6. Human MPs bearing the indicated RAGE alleles were cultured in the presence of either no mediator or EN-RAGE for 14 hrs. Supernatants were retrieved and levels of TNF-alpha (e) and IL-6 (f) determined by ELISA. The mean \pm SD is shown.

15

5G-5H. MMP-9 activity. Human MPs bearing the indicated RAGE alleles were cultured in the presence of either no mediator or EN-RAGE for 14 hrs. Supernatants were retrieved and subjected to zymography to assess levels of activated MMP-9. In b, d, and h, bands from n=5 G82G and n=5 G82S subjects were scanned into a densitometer and band density was quantified. Representative experiments are shown. The mean \pm SD is shown. In b,d,e,f and h, * indicates $p < 0.01$ versus baseline.

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Detailed Description of the Invention

Definitions

5 "Antibody" shall include, by way of example, both naturally occurring and non-naturally occurring antibodies. Specifically, this term includes polyclonal and monoclonal antibodies, and antigen-binding fragments thereof. Furthermore, this term includes chimeric
10 antibodies and wholly synthetic antibodies, and antigen-binding fragments thereof.

"Anti-sense nucleic acid" shall mean any nucleic acid which, when introduced into a cell, specifically
15 hybridizes to at least a portion of an mRNA in the cell encoding a protein ("target protein") whose expression is to be inhibited, and thereby inhibits the target protein's expression.

20 "Catalytic nucleic acid" shall mean a nucleic acid that specifically recognizes a distinct substrate and catalyzes the chemical modification of this substrate. Catalytic nucleic acids include, for example, DNazymes and ribozymes.

25

"DNAzyme" shall mean a catalytic nucleic acid that is DNA or whose catalytic component is DNA, and which specifically recognizes and cleaves a distinct target nucleic acid sequence, which can be either DNA or RNA.
30 Each DNAzyme has a catalytic component (also referred to as a "catalytic domain") and a target sequence-binding component consisting of two binding domains, one on either side of the catalytic domain.

"Expressible nucleic acid" shall mean a nucleic acid encoding a nucleic acid of interest and/or a protein of interest, which nucleic acid is an expression vector, plasmid or other construct which, when placed in a cell, permits the expression of the nucleic acid or protein of interest. Expression vectors and plasmids are well known in the art.

(
As used herein, "inflammatory disorder" includes, without limitation, any disorder possessing an inflammatory component. Examples of inflammatory disorders include autoimmune disorders generally, rheumatoid arthritis, amyotrophic lateral sclerosis, vascular disorders (e.g. atherosclerosis or transplant vasculopathy), and diabetes.

As used herein, "inhibit," when used in connection with the binding between RAGE and/or RAGE G82S with a ligand thereof, shall mean to reduce such binding. In one embodiment, "inhibit" shall mean to eliminate such binding.

"Inhibiting" the onset of a disorder shall mean either lessening the likelihood of the disorder's onset, or preventing the onset of the disorder entirely. In the preferred embodiment, inhibiting the onset of a disorder means preventing its onset entirely.

"Nucleic acid" shall mean any nucleic acid molecule, including, without limitation, DNA, RNA and hybrids thereof. The nucleic acid bases that form nucleic acid molecules can be the bases A, C, G, T and U, as well as derivatives thereof. Derivatives of these bases are well known in the art, and are exemplified in PCR

Systems, Reagents and Consumables (Perkin Elmer Catalogue 1996-1997, Roche Molecular Systems, Inc., Branchburg, New Jersey, USA).

5 "Ribozyme" shall mean a catalytic nucleic acid molecule which is RNA or whose catalytic component is RNA, and which specifically recognizes and cleaves a distinct target nucleic acid sequence, which can be either DNA or RNA. Each ribozyme has a catalytic component (also
10 referred to as a "catalytic domain") and a target sequence-binding component consisting of two binding domains, one on either side of the catalytic domain.

"Specifically hybridize" to a nucleic acid shall mean,
15 with respect to a first nucleic acid, that the first nucleic acid hybridizes to a second nucleic acid with greater affinity than to any other nucleic acid.

"Specifically inhibit" the expression of a protein
20 shall mean to inhibit that protein's expression (a) more than the expression of any other protein, or (b) more than the expression of all but 10 or fewer other proteins.

25 "Subject" shall mean any animal, such as a human, non-human primate, mouse, rat, guinea pig or rabbit.

"Treating" a disorder shall mean slowing, stopping or reversing the disorder's progression. In the preferred
30 embodiment, treating a disorder means reversing the disorder's progression, ideally to the point of eliminating the disorder itself. As used herein,

ameliorating a disorder and treating a disorder are equivalent.

Embodiments of the Invention

5

This invention provides methods for inhibiting binding between RAGE G82S with a ligand thereof, and for inhibiting the onset of and treating inflammatory disorders such as rheumatoid arthritis. This invention
10 is based on the surprising discovery of a correlation between suppressing rheumatoid arthritis and blocking RAGE and/or RAGE G82S function.

Specifically, this invention provides a method for
15 inhibiting binding between RAGE and a ligand thereof comprising contacting the RAGE with an agent comprising (i) soluble RAGE G82S, (ii) a ligand-binding portion of soluble RAGE G82S, or (iii) an antibody directed to RAGE G82S. This invention also provides a method for
20 inhibiting binding between RAGE G82S and a ligand thereof comprising contacting the RAGE G82S with an agent comprising (i) soluble RAGE G82S, (ii) a ligand-binding portion of soluble RAGE G82S, or (iii) an antibody directed to RAGE G82S.

25

In one embodiment of the instant methods, the agent comprises soluble RAGE G82S. In another embodiment, the agent comprises a ligand-binding portion of soluble RAGE G82S. The ligand-binding portion can include, for
30 example, amino acid residues 1-112 of RAGE G82S, and the V-domain. In a further embodiment, the agent comprises an antibody to RAGE G82S. In yet a further embodiment, the agent comprises a small molecule such as an organic molecule.

This invention also provides a method for treating an inflammatory disorder in a subject comprising administering to the subject a therapeutically effective amount of an agent that inhibits binding between RAGE G82S and a ligand thereof. This invention also provides a method for inhibiting the onset of an inflammatory disorder in a subject comprising administering to the subject a prophylactically effective amount of an agent that inhibits binding between RAGE G82S and a ligand thereof.

In the preferred embodiment of the instant methods, the subject is human. In one embodiment the inflammatory disorder is rheumatoid arthritis. In one embodiment of the instant methods, the agent comprises soluble RAGE G82S. In another embodiment, the agent comprises a ligand-binding portion of soluble RAGE G82S. In a further embodiment, the agent comprises an antibody to RAGE G82S. In yet a further embodiment, the agent comprises a small molecule such as an organic molecule.

Determining a therapeutically or prophylactically effective amount of agent can be done based on animal data using routine computational methods. In one embodiment, the therapeutically or prophylactically effective amount contains between about 1ng and about 1g of nucleic acid or protein, as applicable. In another embodiment, the effective amount contains between about 10ng and about 100mg of nucleic acid or protein, as applicable. In a further embodiment, the effective amount contains between about 100ng and about 10mg of the nucleic acid or protein, as applicable. In a yet a further embodiment, the effective amount

contains between about 1 μ g and about 1mg of the nucleic acid or protein, as applicable. In a yet a further embodiment, the effective amount contains between about 10 μ g and about 100 μ g of the nucleic acid or protein, as applicable. In a yet a further embodiment, the effective amount contains between about 100 μ g and about 10mg of the nucleic acid or protein, as applicable.

In this invention, administering agents can be effected or performed using any of the various methods and delivery systems known to those skilled in the art. The administering can be performed, for example, intravenously, orally, via implant, transmucosally, transdermally, intramuscularly, and subcutaneously. The following delivery systems, which employ a number of routinely used pharmaceutical carriers, are only representative of the many embodiments envisioned for administering the instant compositions.

Injectable drug delivery systems include solutions, suspensions, gels, microspheres and polymeric injectables, and can comprise excipients such as solubility-altering agents (e.g., ethanol, propylene glycol and sucrose) and polymers (e.g., polycaprylactones and PLGA's). Implantable systems include rods and discs, and can contain excipients such as PLGA and polycaprylactone.

Oral delivery systems include tablets and capsules. These can contain excipients such as binders (e.g., hydroxypropylmethylcellulose, polyvinyl pyrrolidone, other cellulosic materials and starch), diluents (e.g., lactose and other sugars, starch, dicalcium phosphate and cellulosic materials), disintegrating agents (e.g.,

starch polymers and cellulosic materials) and lubricating agents (e.g., stearates and talc).

Transmucosal delivery systems include patches, tablets, 5 suppositories, pessaries, gels and creams, and can contain excipients such as solubilizers and enhancers (e.g., propylene glycol, bile salts and amino acids), and other vehicles (e.g., polyethylene glycol, fatty acid esters and derivatives, and hydrophilic polymers 10 such as hydroxypropylmethylcellulose and hyaluronic acid).

Dermal delivery systems include, for example, aqueous and nonaqueous gels, creams, multiple emulsions, 15 microemulsions, liposomes, ointments, aqueous and nonaqueous solutions, lotions, aerosols, hydrocarbon bases and powders, and can contain excipients such as solubilizers, permeation enhancers (e.g., fatty acids, fatty acid esters, fatty alcohols and amino acids), and 20 hydrophilic polymers (e.g., polycarbophil and polyvinylpyrrolidone). In one embodiment, the pharmaceutically acceptable carrier is a liposome or a transdermal enhancer.

25 Solutions, suspensions and powders for reconstitutable delivery systems include vehicles such as suspending agents (e.g., gums, xanthans, cellulose and sugars), humectants (e.g., sorbitol), solubilizers (e.g., ethanol, water, PEG and propylene glycol), surfactants 30 (e.g., sodium lauryl sulfate, Spans, Tweens, and cetyl pyridine), preservatives and antioxidants (e.g., parabens, vitamins E and C, and ascorbic acid), anti-caking agents, coating agents, and chelating agents (e.g., EDTA).

In one embodiment of this invention, the delivery system used comprises more than water alone, or more than buffer alone.

5

This invention further provides a method for inhibiting the onset of an inflammatory disorder in a subject comprising administering to the subject a prophylactically effective amount of a nucleic acid
10 which specifically inhibits the expression of RAGE G82S in the subject's cells expressing same. This invention further provides a method for treating a subject afflicted with an inflammatory disorder comprising administering to the subject a therapeutically
15 effective amount of a nucleic acid which specifically inhibits the expression of RAGE G82S in the subject's cells expressing same.

In the preferred embodiment the subject is human. In a
20 one embodiment the inflammatory disorder is rheumatoid arthritis. In one embodiment the nucleic acid is DNA. In another embodiment the nucleic acid is RNA. In a further embodiment the nucleic acid is an anti-sense nucleic acid that specifically hybridizes to mRNA
25 encoding RAGE G82S. In yet a further embodiment the nucleic acid is a catalytic nucleic acid that cleaves mRNA encoding RAGE G82S. In yet a further embodiment the nucleic acid is an expressible nucleic acid encoding an anti-sense nucleic acid that specifically
30 hybridizes to mRNA encoding RAGE G82S. In yet a further embodiment the nucleic acid is an expressible nucleic acid encoding a catalytic nucleic acid that cleaves mRNA encoding RAGE G82S. In one embodiment the subject's cells in which the amount of RAGE G82S is to

be reduced are cells which mediate an inflammatory reaction.

This invention further provides soluble RAGE G82S, a
5 ligand-binding portion thereof, nucleic acids encoding same, and anti-RAGE G82S antibodies. The sequence of human RAGE is known (10, 11, and 61), as are ligand-binding portions thereof (62, 63).

10 This invention further provides a composition comprising a pharmaceutically acceptable carrier and (i) soluble RAGE G82S, (ii) a ligand-binding portion of soluble RAGE G82S, and/or (iii) a nucleic acid which specifically inhibits the expression of RAGE G82S in a
15 cell expressing same. Making and using nucleic acids to selectively inhibit protein expression is known, as exemplified, for example, in patents relating to gene therapy (64-68), anti-sense therapy (69 and 70) and dsRNA use (71).

20 Finally, this invention provides an article of manufacture comprising a packaging material having therein (i) an agent that inhibits binding between RAGE G82S and a ligand thereof and/or (ii) a nucleic acid
25 which specifically inhibits the expression of RAGE G82S in a cell expressing same, wherein the packaging material has affixed thereto a label indicating a use for the agent and/or nucleic acid for treating an inflammatory disorder in a subject.

30 In the preferred embodiment of the instant article of manufacture, the subject is human. In one embodiment of the invention, the disorder is rheumatoid arthritis and the agent comprises soluble RAGE G82S, a ligand-

binding portion of soluble RAGE G82S, or an antibody directed to RAGE G82S. In one embodiment of the instant article of manufacture, the agent comprises soluble RAGE G82S. In another embodiment, the agent
5 comprises a ligand-binding portion of soluble RAGE G82S. In a further embodiment, the agent comprises an antibody to RAGE G82S. In yet a further embodiment, the agent comprises a small molecule such as an organic molecule.

10

This invention is illustrated in the Experimental Details section which follows. This section is set forth to aid in an understanding of the invention but is not intended to, and should not be construed to,
15 limit in any way the invention set forth in the claims which follow.

Experimental Details

Murine model of bovine collagen type II-induced arthritis

5

Induction of arthritis. Male DBA/1 mice were purchased from the Jackson Laboratories (Bar Harbor, ME). On day 0, mice weighing 20-30 gms were injected intradermally at the base of the tail with bovine type II collagen, 10 200 µg (Sigma), dissolved in acetic acid (0.01M) and emulsified in complete Freund's adjuvant (ICN Biochemcials, Costa Mesa, CA). Three weeks later (day 22), mice were challenged by injection of bovine collagen type II (200 µg) in incomplete Freund's 15 adjuvant (15-17). Beginning three weeks after immunization (day 22), mice were treated with murine soluble RAGE (1) (100 µg per day by intraperitoneal route). Vehicle treatment consisted of murine serum albumin (Sigma), 100 µg per day by intraperitoneal 20 injection, or equal volumes of phosphate-buffered saline (PBS). Treatment was continued daily until sacrifice. All murine studies were performed in accordance with the policies of Columbia University's Institutional Animal Care and Use Committee.

25

Determination of anti-collagen IgG. ELISA was performed to determine levels of IgG to bovine type II collagen in dba/1 mice. Plastic Maxisorp 96-well plates (Nunc, Naperville, ILL) were coated with bovine 30 type II collagen (5 µg/ml) in bicarbonate/carbonate buffer; pH 9.3 for 16 hrs at 4°C. Unbound sites were blocked with PBS containing BSA (1%) for 8 hours at 4°C. Wells were washed five times in PBS and then incubated for 16 hrs in samples of mouse plasma 35 (dilution, 1:50) in PBS at 4°C. Wells were washed five

times with PBS and incubated with PBS containing 1% BSA and horse radish peroxidase-labelled anti-mouse IgG (Sigma; 1:500 dilution) for two hours at room temperature. Wells were washed with PBS five times and developed using o-phenylenediamine dihydrochloride (Sigma). A standard curve was prepared using serial dilutions from a mouse antibody cocktail against bovine type II collagen (Chemicon, Temecula, CA).

10 *Assessment of arthritis.* Arthritis was evaluated by clinical and histologic scoring by observers blinded to the experimental conditions. Clinical severity of arthritis in each of four limbs was assessed by the following scale (19): 0=normal; 1=slight erythema and swelling; 2=pronounced edematous swelling; and 3=joint deformity with ankylosis, resulting in a maximum score of 12 per animal. Histologic scoring of arthritis and cartilage and bone destruction was performed on hematoxylin and eosin-stained sections of extremity joints by the following scale (20): (i) synovial lesions: 0, no lesions; 1: mild effect; 2: moderate effect/proliferation; 3: severe lesions with destruction; (ii) cartilage destruction: 0, none; 1, mild; 2, moderate; 3, severe destruction with loss or complete fragmentation of cartilage; and (iii) bone destruction: 0, none; 1, mild destruction of subchondral bone; 2, moderate destruction; and 3, severe destruction with loss of large areas of bone. In other studies, six weeks after initial sensitization, bovine type II collagen (10 µg) was injected into the ear of each mouse. 18 hrs later, thickness of the ear was assessed using calipers by an observer blinded to the experimental conditions.

Immunoblotting and ELISA. Mice were sacrificed 3 or 6 weeks after challenge. The stifle joint was removed and homogenized in tris-buffered saline containing protease inhibitors (Complete Protease Inhibitor, 5 Boehringer-Mannheim, Indianapolis, IN). SDS-PAGE and immunoblotting were performed on extracts of stifle joint tissue using the following antibodies: anti-RAGE IgG and anti-S100/calgranulin IgG as previously described (1) (4.7 and 2.0 µg/ml, respectively). ELISA 10 for TNF-alpha and IL-6 was performed using kits from R&D Systems (Minneapolis, MN).

Zymography. Zymography for detection of MMP-3, 9 and 13 activity was performed using gelatin or casein-laden 15 gels from Novex/Invitrogen (Carlsbad, CA).

Chinese Hamster Ovary (CHO) cell studies

Chinese hamster ovary (CHO) cells were obtained from 20 ATCC (Rockville, MD) and cultured in F12 medium containing fetal bovine serum (10%) (Life Technologies). To generate the 82S allele, the cDNA encoding human RAGE (10) was cloned using the TOPO TA cloning system into pCR2.1TOPO vector for mutagenesis 25 (Invitrogen, Carlsbad, CA). Site-directed mutagenesis to insert the (-82G-) to (-82S-) change was performed using the GeneEditor In Vitro SDM System (Promega, Madison, WI). Sequencing was performed using an ABI310 automated DNA sequencer (Perkin Elmer Biosystems, 30 Foster City, CA) to confirm the inserted sequence changes and to ensure that no other mutations were created. RAGE 82G and 82S cDNAs were excised from pCR2.1TOPO using EcoR I and subcloned into the pcDNA3.1 expression vector (Invitrogen). Cells were transfected

with plasmid DNA using lipofectamine (Life Technologies) encoding the following: pCDNA3.1 containing full-length RAGE cDNA (-82G-), pCDNA3.1 containing RAGE cDNA (-82S-), or pCDNA3.1 containing no insert (mock). 24 hrs after transfection, selection was begun using G418 (1 mg/ml) (Life Technologies). RAGE expression was assessed by immunoblotting in stably-transfected cells after 6 weeks. Cells were incubated with EN-RAGE (Extracellular Newly-Identified RAGE binding protein), a prototypic S100/calgranulin molecule previously-identified to be a signal transduction ligand for RAGE (1), or the indicated mediators. Assessment of activation of phosphorylated MEK 1/2 and p44/p42 MAP kinase by immunoblotting or nuclear translocation of NF-kB was performed. All reagents were tested for endotoxin content using the limulus amebocyte assay (Sigma). Any endotoxin identified in these assays was removed by chromatography of the protein solutions onto Detox-igel columns (Pierce, Arlington Hts, ILL). The material was re-tested in the limulus amebocyte assay for the absence of detectable endotoxin.

Radioligand binding assays. Purified EN-RAGE was radiolabelled using ¹²⁵I and Iodobeads (Pierce, Arlington Heights, ILL) to a specific activity of approximately 5,000 cpm/ng. Radioligand binding assays were performed in 96-well tissue culture dishes containing the indicated transfected CHO cells in the presence of the indicated concentration of radiolabelled EN-RAGE ± an 50-fold molar excess of unlabelled EN-RAGE in PBS containing calcium/magnesium and BSA, 0.2%, for 3 hrs at 37°C. Wells were washed rapidly with washing buffer (PBS containing Tween 20

(0.05%)). Elution of bound material was performed in a solution containing heparin, 1 mg/ml. The aspirate was counted in a gamma counter (LKB, Gaithersburg, MD). Equilibrium binding data were analyzed according to the equation of Klotz and Hunston (57): $B = nKA / 1 + KA$, where B=specifically bound ligand (total binding, wells incubated with tracer alone, minus nonspecific binding, wells incubated with tracer in the presence of excess unlabelled material), n=sites/cell, K=the dissociation constant, and A=free ligand concentration) using nonlinear least-squares analysis (Prism; San Diego, CA). Where indicated, pretreatment with antibodies (70 µg/ml), or human soluble RAGE (50-fold molar excess), was performed.

15 *Activation of pMEK 1/2 and p44/p42 MAP kinases.* CHO cells were incubated with EN-RAGE, 10 µg/ml, for one hr. Cells were lysed in lysis buffer (New England Biolabs, Beverly, MA) and lysate was subjected to centrifugation. Protein concentration of the supernatant was determined by Bio-Rad assay (Bio-Rad, Hercules, CA). Equal amounts of protein were subjected to SDS-PAGE (Novex/Invitrogen). Contents of the gels were transferred to nitrocellulose and immunoblotting performed using anti-phosphorylated/total p44/p42 MAP kinase and MEK 1/2 (New England Biolabs) (1 µg/ml).

Electrophoretic mobility shift assay. Nuclear extracts were prepared using the NE kit (Pierce) and EMSA performed employing consensus ³²P-labelled probe for NF-kB as described (1). Where indicated, cells were treated with either nonimmune/anti-RAGE IgG, or soluble RAGE (1,21).

Peripheral blood-derived mononuclear phagocyte (MPs) studies

Cellular isolation. Venous blood was obtained from
5 healthy volunteers (30 ml) bearing RAGE 82G or 82G/82S
in accordance with the policies of Columbia
University's Institutional Review Board. Due to the
rare numbers of subjects homozygous for the 82S allele,
the studies were confined to those bearing solely 82G
10 or heterozygous 82G/82S. Mononuclear cells were
isolated using Histopaque 1077 (Sigma, St. Louis, MO)
and cultured on plastic dishes for 3 hrs at 37°C.
Nonadherent cells were removed by washing in PBS.
Adherent cells were removed by incubation with EDTA (2
15 mM) for 15 mins at 37°C. Cells (>95% MPs as assessed
by immunostaining with anti-CD68 IgG) were seeded in
tissue culture-coated wells for study.

Activation of phosphorylated MEK 1/2 and p44/p42 MAP
20 *kinases and generation of IL-6 and TNF-alpha.* MPs
were seeded onto the wells of 24-well tissue culture
plates at a density of 5×10^5 cells per well. Cells were
stimulated with either BSA or EN-RAGE (10 µg/ml), and
immunoblotting for detection of phosphorylated/total
25 MEK 1/2 and p44/p42 MAP kinases performed as above.
Supernatant was assayed for IL-6 and TNF-alpha using
ELISA kits from R&D systems (Minneapolis, MN).

Patient population and detection of 82S polymorphism

30 RA patients used for association studies meet the
criteria of the American College of Rheumatology (58)
and were taken from patient populations collected
previously at the Medical College of Virginia, Virginia
35 Commonwealth University (31). The following primers

were synthesized for detection of the glycine82serine (G82S) polymorphism of the RAGE gene (8): sense primer: 5' GTAAGCGGGGCTCCTGTTGCA-3' and the antisense primer: 5' GGCCAAGGCTGGGGTTGAAGG-3'. Whole blood (20 µl) was obtained from human volunteers in accordance with the standards and policies of the Institutional Review Boards of the institutions. Genomic DNA was prepared using a kit from Qiagen (Valencia, CA); 10 ng was amplified using Taq DNA polymerase (Life Technologies, Grand Island, NY) in a final volume of 25 µl and PCR performed as follows: 94°C for 30 secs, 62°C for 45 secs, and 72°C for 60 secs for a total of 35 cycles. PCR product (25 µl) was digested with Alu I (Life Technologies), 3U for 16 hr at 37°C, followed by electrophoresis on 2% agarose gels.

Statistical analysis

Statistical comparisons were determined using one-way analysis of variance (ANOVA); where indicated, individual comparisons were performed using students' t-test. For cell culture studies, immunoblotting, zymography and electrophoretic mobility shift assays were performed at least three times. Statistical analyses were performed across each of the experiments by scanning gels/autoradiograms into a densitometer. Arbitrary units of intensity (Image Quant/Molecular Dynamics (Foster City, CA)) were then standardized using untreated cells as control, and assigned a value of "1". Similarly, in animal studies, an arbitrary value of "1" was assigned to bands (immunoblotting or zymography) in assays from animals without collagen-induced arthritis). Statistical analyses to identify the incidence of G82S in human subjects were performed

as previously described (59-60). $P < 0.05$ was considered statistically significant.

Results

5

RAGE and Murine Collagen-Induced Arthritis

As a first test of the hypothesis that RAGE contributed to inflammatory and tissue-destructive mechanisms in arthritis, arthritis was induced in DBA/1 mice by sensitization and challenge with bovine type II collagen, the predominant protein of articular cartilage (15-17). Bovine type II collagen was emulsified in complete Freund's adjuvant and injected intradermally at the base of the tail (day 0). Three weeks later, mice were challenged with bovine type II collagen (day 22). The contribution of RAGE to the pathogenesis of arthritis was studied by treating animals with soluble (s) RAGE (1,18), the extracellular ligand-binding domain of the receptor, 100 μ g per day, beginning three weeks after initial immunization (day 22). In previous studies, blockade of RAGE at this dose effected the greatest decrease in the proinflammatory phenotype in a murine model of delayed-type hypersensitivity (1).

The relevance of RAGE-ligand interaction to murine collagen-induced arthritis was underscored by the increased expression of RAGE and S100/calgranulins in joint tissue. Six weeks after immunization, RAGE and S100/calgranulin expression was increased in joints from mice with arthritis compared with controls, especially within the proliferating synovium (RAGE, fig. 1e&d; S100/calgranulin, fig. 1h&g, respectively).

By immunoblotting, levels of RAGE antigen were enhanced ~2.2-fold in arthritis versus control joints (fig. 1m). Although levels of S100/calgranulins were quite low in joint tissue of control DBA/1 mice without collagen-induced arthritis, expression of these proinflammatory mediators was induced in murine serum albumin-treated mice immunized/challenged with bovine type II collagen (fig. 1n). In mice subjected to blockade of RAGE with sRAGE, levels of RAGE and S100/calgranulin antigens by immunoblotting were significantly reduced compared to mice treated with vehicle (fig. 1m&n, respectively).

In parallel with reduced expression of RAGE and S100/calgranulins in sRAGE-treated joints, clinical indices of inflammation (18) were consistently lower in mice treated with sRAGE vs PBS or murine serum albumin (fig. 2a). In addition, histologic scoring (19) revealed marked decreases in synovial hyperplasia/hypertrophy in the joints of sRAGE-treated mice (fig. 2b). Further, histologic indices of cartilage and bone destruction (20) were significantly decreased in sRAGE-treated mice compared with those animals receiving PBS or murine serum albumin (fig. 2b). To determine if blockade of RAGE suppressed immune/inflammatory responses to bovine type II collagen at extra-articular sites, mice receiving vehicle or sRAGE were injected with bovine type II collagen (10 µg) into ear tissue six weeks after immunization (day 42). Although baseline ear thickness was identical in all groups, 18 hrs after injection of collagen, mice receiving murine serum albumin or PBS revealed significant increases in ear thickness compared with those mice treated with sRAGE (fig. 2c). Levels of serum total IgG developed against bovine type

II collagen were unchanged in the presence of sRAGE vs PBS or murine serum albumin, thereby indicating that blockade of RAGE did not suppress primary immunization (fig. 2d).

5

To dissect the molecular mechanisms underlying the protection afforded by RAGE blockade, levels of cytokines and matrix metalloproteinases (MMPs) in the tissues were assessed. ELISA on extracts prepared from
10 stifle joints revealed a significant reduction in levels of TNF-alpha (~4.3-fold) and IL-6 (~3.1-fold) from mice immunized with bovine type II collagen treated with sRAGE vs murine serum albumin (fig. 3a&b, respectively). Similarly, levels of plasma TNF-alpha
15 and IL-6 antigens were significantly reduced, ~3.5- and 3.6-fold, in sRAGE-treated mice compared with those animals receiving murine serum albumin (fig. 3c&d, respectively).

20 As induction of TNF-alpha and other inflammatory cytokines sets in motion events leading to activation of latent/proenzyme MMPs (21), activity of MMPs 3, 9 and 13 in joint tissue was assessed. Compared with control joints, tissue retrieved from murine serum albumin-
25 treated mice undergoing the collagen-induced arthritis protocol revealed an ~2.5-fold, ~4.0-fold and ~5.3-fold increase in activity of MMPs 3, 9 and 13, respectively, by zymography (fig. 3e&f). That activation of RAGE was critical in this process was demonstrated by the
30 significant reduction in activation of these MMPs in joint tissue of sRAGE-treated mice (fig. 3e&f, respectively).

Taken together, these observations suggested that interaction of S100/calgranulins and RAGE, both enriched in joint tissues in arthritis, activates proinflammatory and tissue-destructive mediators, thereby supporting roles for these molecules in the evolution of joint destruction.

RAGE 82S Allele Enhances Binding/Signalling/Gene Expression by S100/calgranulins

10

Polymorphisms within the RAGE gene may influence these effects by altering proinflammatory pathways. To study this, whether the glycine to serine polymorphism (G82S) within the ligand-binding domain of RAGE displayed enhanced ligand affinity and activation of signal transduction pathways was tested. Stably-transfected Chinese hamster ovary (CHO) cells bearing the two RAGE alleles of interest were prepared, either the common RAGE 82G allele, or the variant RAGE 82S allele. CHO cells provide a convenient model, as they are devoid of detectable RAGE prior to (not shown), or after stable transfection with pcDNA3.1 vector alone (mock) (fig. 4a). Stably-transfected CHO cells were made with pcDNA3.1 containing either RAGE 82G or RAGE 82S (fig. 4a). ELISA studies demonstrated that levels of RAGE antigen were identical in both CHO cells transfected to express RAGE 82G or RAGE 82S (data not shown). Radioligand binding studies showed dose-dependent binding of ^{125}I -EN-RAGE (a prototypic S100/calgranulin) to CHO cells expressing RAGE 82G ($\sim 122 \pm 31$ nM) and 82S receptor ($K_d \sim 77 \pm 21$ nM), though the affinity of binding was greater with the 82S allele ($p=0.008$; fig. 4b). In contrast, CHO cells stably transfected with the empty vector displayed no specific binding of ^{125}I -EN-RAGE (fig. 4b). That the interaction of RAGE-bearing CHO

cells with ^{125}I -EN-RAGE was specific for interaction with RAGE was shown by suppression of specific binding in the presence of excess soluble extracellular domain of the receptor (sRAGE), or anti-RAGE IgG, but not by
5 bovine serum albumin (BSA) or nonimmune IgG (fig. 4c).

These observations led to testing the concept that the 82S RAGE isoform might amplify cellular activation beyond that seen in cells bearing wild-type RAGE.

10 Incubation of mock-transfected CHO cells with a physiologically-relevant concentration of EN-RAGE (4), 10 $\mu\text{g/ml}$, did not significantly increase intensity of the bands corresponding to phosphorylated MEK 1/2 or p44/42 MAP kinases (22-23) (figs. 4d&e, respectively;
15 lanes 1-5). However, exposure of RAGE 82G CHO transfectants to EN-RAGE increased by ~ 2.2 -fold and ~ 1.9 -fold phosphorylated MEK 1/2 and p44/42 MAP kinases, compared with cultures incubated with BSA alone ($p < 0.01$; fig. 4d&e, respectively; lanes 7&6).

20 CHO transfectants bearing RAGE 82S incubated with EN-RAGE displayed ~ 3.6 -fold and ~ 4.1 -fold increase in phosphorylated MEK 1/2 and p44/42 compared with BSA ($p < 0.01$; fig. 4d&e, respectively; lanes 12&11). Thus, compared with cells expressing the common RAGE 82G
25 allele, CHO cells expressing the variant RAGE 82S displayed significantly increased phosphorylation of MEK 1/2 and p44/p42 MAP kinases by ~ 1.6 - and ~ 2.2 -fold ($p < 0.05$; fig. 4d&e, respectively; lanes 12&7). In both RAGE 82G and RAGE 82S-transfected cells, cellular
30 activation by EN-RAGE was due to engagement of RAGE as demonstrated by suppression of phosphorylation of MEK 1/2 and p44/p42 by excess sRAGE (fig. 4d&e, respectively; lanes 8&13), or anti-RAGE IgG (fig. 4d&e, respectively, lanes 9&14). Nonimmune IgG was without

effect (fig. 4d&e, respectively; lanes 10&15). In each case, respective levels of total MEK 1/2 and p44/p42 MAP kinases were identical (data not shown).

5 To further support the hypothesis that RAGE 82S enhanced activation of key proinflammatory signal transduction pathways, nuclear translocation of NF- κ B in CHO transfectants exposed to EN-RAGE were assessed. Electrophoretic mobility shift assays (EMSA) using 32 P-
10 labelled consensus NF- κ B probe and nuclear extracts from mock-transfected CHO cells showed no increase in intensity of the gel shift band after cultures were exposed to EN-RAGE (fig. 4f, lane 2). In CHO transfectants expressing RAGE 82G, there was a
15 prominent ~2.4-fold increase in intensity in nuclear binding activity following incubation of cultures with EN-RAGE compared to BSA ($p < 0.05$; fig. 4f, lanes 7&6, respectively). NF- κ B activation was even more striking when RAGE 82S was substituted for RAGE 82G. RAGE 82S
20 CHO transfectants displayed ~5.1-fold increased intensity of the gel shift band consequent to the presence of EN-RAGE, compared to incubation with BSA ($p < 0.001$; fig. 4f, lanes 12&11, respectively). Thus, RAGE-mediated NF- κ B activation due to EN-RAGE was
25 significantly enhanced by ~2.1-fold comparing RAGE 82S to 82G; $p < 0.01$. That activation of NF- κ B in transfected CHO cells by EN-RAGE resulted from ligation of common or variant RAGE was confirmed by its suppression in the presence of sRAGE (fig. 4f, lanes
30 8&13, respectively), or anti-RAGE IgG (fig. 4f, lanes 9&14, respectively), but not by nonimmune IgG (fig. 4f, lanes 10&15, respectively).

A critical test of the concept that S100/calgranulin-RAGE 82S interaction might enhance proinflammatory effects was whether mononuclear phagocytes (MPs) retrieved from human subjects bearing the variant RAGE 82S allele displayed heightened activation of signal transduction molecules in the presence of ligand. Due to the rare number of subjects homozygous for 82S in our study populations, studies were limited to those individuals homozygous for the common RAGE allele 82G, or those subjects heterozygous for 82G and 82S alleles. ELISA revealed that basal levels of RAGE antigen did not differ between MPs bearing 82G or 82G/82S (data not shown). Signalling was compared in MPs from subjects bearing RAGE 82G and RAGE 82G/82S by assessing activation of MEK 1/2 and p44/p42 MAP kinases. In the presence of EN-RAGE, MPs isolated from individuals bearing RAGE 82G/82S displayed an ~4.5- and 4.3-fold increase in phosphorylated MEK 1/2 and p44 and p42 MAP kinases, respectively, compared with unstimulated cells ($p < 0.01$; fig. 5a,b,c&d). However, MPs bearing the common RAGE 82G allele exposed to EN-RAGE revealed a significant, although smaller (~2.6- and 2.3-fold) increase in activation of MEK 1/2 and p44/p42 MAP kinases, respectively ($p < 0.01$; fig. 5a,b,c&d). Importantly, EN-RAGE-triggered activation of MEK 1/2 and p44/p42 MAP kinases was significantly increased in heterozygous RAGE 82G/82S-bearing MPs vs those MPs expressing RAGE 82G ($p < 0.05$; fig. 5b&d).

To assess the functional consequences of enhanced binding and activation of signal transduction molecules stimulated upon ligation of RAGE by EN-RAGE, production of key inflammatory and tissue-degradative mediators linked to RA (24-26) by MPs bearing common or variant

RAGE alleles were examined. Exposure of RAGE 82G-bearing MPs to EN-RAGE caused a significant increase in generation of TNF-alpha detected in culture supernatant compared with quiescent cultures (470 ± 53 vs 15 ± 2 pg/ml; $p < 0.001$) (fig. 5e). However, upon incubation of human MPs bearing RAGE 82G/82S, an even greater increase in elaborated TNF-alpha was observed in culture supernatants compared to basal levels (850 ± 64 vs 17 ± 2 pg/ml; $p < 0.01$) (fig. 5e). Importantly, although basal levels of TNF-alpha did not differ between RAGE 82G and 82G/82S-bearing MPs, levels of TNF-alpha in EN-RAGE-stimulated MPs were increased ~1.8-fold in the presence of RAGE 82G/82S compared with RAGE 82G ($p < 0.01$; fig. 5e). Similarly, RAGE 82G-bearing MPs exposed to EN-RAGE displayed a small, but significant increase in generation of IL-6 compared with basal expression (55 ± 7 vs 15 ± 2 pg/ml; $p < 0.01$) (fig. 5f). However, MPs bearing RAGE 82G/82S revealed augmented generation of IL-6 upon incubation with EN-RAGE compared with unstimulated controls (160 ± 14 vs 12 ± 2 pg/ml; $p < 0.01$) (fig. 5f). EN-RAGE-stimulated MPs bearing RAGE 82G/82S generated increased amounts of IL-6 compared with cells from subjects bearing RAGE 82G (~2.9-fold) ($p < 0.01$; fig. 5f).

25

A central means by which structural elements of joints are degraded in unchecked RA is by generation of MMPs, such as MMP-9 (27-30). It was hypothesized that RAGE-mediated MP activation would augment generation of MMP activity on cells bearing RAGE 82G/82S versus RAGE 82G. Activity of MMP-9 in MPs retrieved from subjects bearing the common or variant RAGE alleles was tested. MPs from subjects bearing RAGE 82G displayed an ~2.3-fold increase in MMP-9 activity in the presence of EN-

RAGE compared with basal expression ($p < 0.01$; fig. 5g&h). However, MPs isolated from individuals bearing RAGE 82G/82S demonstrated an ~4.3-fold increase in EN-RAGE-mediated MMP-9 activity compared with baseline (p<0.01; fig. 5g&h). Although basal levels of MMP-9 activity did not differ among 82G- or 82G/82S-bearing MPs, the extent of EN-RAGE-mediated enhanced MMP-9 activity was significantly enhanced, ~1.9-fold, in MPs bearing RAGE 82G/82S vs RAGE 82G ($p < 0.05$; fig. 5g&h).

10

Taken together, the studies in DBA/1 mice suggest that activation of RAGE amplifies cytokine generation and activation of MMPs in joint tissue primed by immunization with bovine type II collagen. These considerations, together with the findings in vitro suggesting that S100/calgranulin-RAGE interaction modifies cellular responses, especially in the presence of the 82S allele, led to the evaluation of the association of the RAGE 82S allele with RA.

20

RAGE 82S Allele and Association with Rheumatoid Arthritis

Subjects from a previously-reported Caucasian population of RA patients (n=205) and matched controls (n=169) (31) were evaluated. The RAGE 82S allele exhibited an association with RA with relative risk (RR) =2.6, $p < 0.001$ (table 1a). Further analyses were performed to account for the linkage disequilibrium described previously between RAGE 82S and an RA-associated allele, HLA-DRB*0401 (9,32-34). To assess whether the RAGE 82S allele confers risk independent of DRB1*0401, the frequency of the RAGE 82S allele in 0401+ patients and controls was compared. This analysis did not reveal a significant independent effect of RAGE

82S on RA disease susceptibility; $p=0.13$ by Fisher's exact test (table 1b). However, DRB1*0401 continued to exhibit a significant association with RA in the absence of the RAGE 82S allele (table 1c).

5

Discussion

S100/calgranulin molecules have been extensively linked to immune/inflammatory diseases. Although their
10 intracellular functions are relatively well-defined, such as calcium-dependent chemotaxis, degranulation and phagocytosis by activated granulocytes (5-6,35), an increasingly recognized view is that these molecules may exhibit distinct functions in the extracellular
15 milieu (35). These concepts are consistent with the observations that the degree of elevation of S100/calgranulins in inflamed foci may mirror the extent of disease activity. For example, in addition to their association with disease severity in RA (4),
20 S100/calgranulins have been linked to disease progression in inflammatory bowel disease (36,37). Furthermore, strongly increased expression of a member of this family of molecules, psoriasin, has been demonstrated in psoriatic lesions compared with
25 adjacent unaffected skin in human subjects (38), thus further supporting the premise that S100/calgranulins may not simply be "innocent bystanders," but, rather, by virtue of their engagement of RAGE (1), active participants in the host inflammatory response. One
30 can speculate that triggered by interaction with accumulating S100/calgranulins, activation of RAGE engages key signalling pathways linked to proinflammatory responses, such as MAP kinases and NF- κ B, thereby amplifying expression of molecules linked

to chronic cellular perturbation and tissue destruction. This hypothesis appears to be supported by the studies in DBA/1 mice immunized/challenged with bovine type II collagen, as activation of RAGE
5 mediated, at least in part, increased generation of inflammatory cytokines and activation of MMPs. Indeed, blockade of the receptor was associated with decreased clinical and histologic indices of inflammation and bone/cartilage destruction in this model.

10

Polymorphic differences in the RAGE gene may lead to varied effects of RAGE in inflammatory settings. This led to the investigation of the effects of the G82S polymorphism in the binding domain of RAGE on receptor
15 function and on the inflammatory response. In transfected CHO cells, increased ligand affinity was found to be conferred by the 82S isoform, which, upon further analysis, demonstrated upregulation of intracellular signalling pathways linked to modulation
20 of proinflammatory genes upon interaction with S100/calgranulin. Using isolated MPs from human subjects, significant differences in inflammatory responses between cells isolated from homozygous wild-type G82G and heterozygous G82S subjects were
25 demonstrated. These effects were quite marked between wild-type and heterozygote subjects, thereby suggesting that the 82S allele alone may impart an even greater effect.

30 Finally, the prevalence of the 82S allele in a population of subjects with RA was investigated. The results demonstrated a statistically-significant increase in prevalence of the 82S allele in subjects with RA. Sub-analysis for the presence/absence of

DRB1*0401 led to the loss of an statistically-significant association of 82S with RA ($p=0.13$ despite odds ratio 3.2). Conversely, DRB1*0401 continued to exhibit a significant association with RA in the
5 absence of the RAGE 82S allele. Clearly, the large genetic susceptibility to RA related to the MHC (39-40) cannot be ascribed solely to the RAGE 82S allele. Due to the relatively low frequency of the 82S allele and low subject numbers in these Caucasian cases and
10 controls after sub-analysis, it remains possible that the RAGE G82S polymorphism is related independently to RA.

The literature is conflicting with respect to a role
15 for DR4 haplotypes in disease severity (41-50). In view of the linkage disequilibrium between DRB1*0401 and the RAGE 82S allele, it will be interesting to specifically test whether DR4 haplotypes containing the RAGE 82S
20 allele have an influence on disease progression and outcome. By analogy with the observations in cultured CHO and human mononuclear cells, one can speculate it may be possible to identify a subset of subjects with RA particularly vulnerable to the sequelae of
25 heightened production/activation of proinflammatory mediators. Specifically, in parallel with outcome measures of clinical severity, such as numbers of bony erosions, the extent of elevation of TNF-alpha, IL-6 and MMP activities in rheumatoid joint fluid and/or RA plasma must be correlated with subject genotype.

30

A general implication of these studies is that the classical DRB1 associations with RA may not completely explain the underlying reasons for the strong linkage and association of the MHC with RA. Indeed, a number

of recent reports indicate that additional genes in the TNF region may influence disease susceptibility, independent of DRB1, and unrelated to DR4 haplotypes (51-52). A full understanding of the role of the MHC in RA and other autoimmune diseases will require a more detailed analysis of this genetic region, since there are many genes within the MHC, in addition to RAGE, which may influence the immune/inflammatory response (53).

10

Lastly, in addition to the 82S variant of the RAGE gene, it will be important to examine the potential role of other variants of RAGE in disease susceptibility and/or severity in immune/inflammatory diseases. In this context, recently identified polymorphisms within the promoter of the gene encoding RAGE may hold the key to unique elements controlling enhanced transcription of the RAGE gene and increased expression of cell surface RAGE (54). As a number of studies have suggested that expression of RAGE is increased at sites of ligand accumulation such as diabetic atherosclerotic lesions, infected periodontium or Alzheimer disease brain (13,18,55), polymorphisms within regulation elements and/or ligand-binding regions may, alone or in combination, orchestrate RAGE's availability and/or functional potential in a given milieu (56).

In conclusion, one can propose that these observations in an animal model of inflammatory arthritis and in S100/calgranulin-stimulated activation of 82S RAGE-bearing cells highlight ligand-RAGE engagement as an amplifier of proinflammatory mechanisms in immune/inflammatory diseases. These findings provide a

compelling rationale to expand the analysis of RAGE polymorphisms and investigate their potential contribution to susceptibility and/or progression of the inflammatory response in diverse diseases with an
5 immune/inflammatory component, such as RA, atherosclerosis and diabetes.

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What is claimed is:

1. A method for inhibiting binding between RAGE and a ligand thereof comprising contacting the RAGE with an agent comprising (i) soluble RAGE G82S, (ii) a ligand-binding portion of soluble RAGE G82S, or (iii) an antibody directed to RAGE G82S.
2. The method of claim 1, wherein the agent comprises soluble RAGE G82S.
3. The method of claim 1, wherein the agent comprises a ligand-binding portion of soluble RAGE G82S.
4. The method of claim 1, wherein the agent comprises an antibody directed to RAGE G82S.
5. A method for inhibiting binding between RAGE G82S and a ligand thereof comprising contacting the RAGE G82S with an agent comprising (i) soluble RAGE G82S, (ii) a ligand-binding portion of soluble RAGE G82S, or (iii) an antibody directed to RAGE G82S.
6. The method of claim 5, wherein the agent comprises soluble RAGE G82S.
7. The method of claim 5, wherein the agent comprises a ligand-binding portion of soluble RAGE G82S.
8. The method of claim 5, wherein the agent comprises an antibody directed to RAGE G82S.

9. A method for treating an inflammatory disorder in a subject comprising administering to the subject a therapeutically effective amount of an agent that inhibits binding between RAGE G82S and a ligand thereof.
10. The method of claim 9, wherein the inflammatory disorder is rheumatoid arthritis.
11. The method of claim 9, wherein the subject is a human.
12. The method of claim 9, wherein the agent comprises (i) soluble RAGE G82S, (ii) a ligand-binding portion of soluble RAGE G82S, or (iii) an antibody directed to RAGE G82S.
13. The method of claim 12, wherein the agent comprises soluble RAGE G82S.
14. The method of claim 12, wherein the agent comprises a ligand-binding portion of soluble RAGE G82S.
15. The method of claim 12, wherein the agent comprises an antibody directed to RAGE G82S.
16. A method for inhibiting the onset of an inflammatory disorder in a subject comprising administering to the subject a prophylactically effective amount of an agent that inhibits binding between RAGE G82S and a ligand thereof.

17. The method of claim 16, wherein the subject is a human.
18. The method of claim 16, wherein the agent
5 comprises (i) soluble RAGE G82S, (ii) a ligand-binding portion of soluble RAGE G82S, or (iii) an antibody directed to RAGE G82S.
19. The method of claim 18, wherein the agent
10 comprises soluble RAGE G82S.
20. The method of claim 18, wherein the agent
comprises a ligand-binding portion of soluble RAGE
G82S.
- 15 21. The method of claim 18, wherein the agent
comprises an antibody directed to RAGE G82S.
22. A method for inhibiting the onset of an
20 inflammatory disorder in a subject comprising
administering to the subject a prophylactically
effective amount of a nucleic acid which
specifically inhibits the expression of RAGE G82S
in the subject's cells expressing same.
- 25 23. A method for treating a subject afflicted with an
inflammatory disorder comprising administering to
the subject a therapeutically effective amount of
a nucleic acid which specifically inhibits the
30 expression of RAGE G82S in the subject's cells
expressing same.
24. The method of claim 22 or 23, wherein the nucleic
acid is DNA.

25. The method of claim 22 or 23, wherein the nucleic acid is RNA.
- 5 26. The method of claim 22 or 23, wherein the nucleic acid is an anti-sense nucleic acid that specifically hybridizes to mRNA encoding RAGE G82S.
- 10 27. The method of claim 22 or 23, wherein the nucleic acid is a catalytic nucleic acid that cleaves mRNA encoding RAGE G82S.
- 15 28. The method of claim 22 or 23, wherein the nucleic acid is an expressible nucleic acid encoding an anti-sense nucleic acid that specifically hybridizes to mRNA encoding RAGE G82S.
- 20 29. The method of claim 22 or 23, wherein the nucleic acid is an expressible nucleic acid encoding a catalytic nucleic acid that cleaves mRNA encoding RAGE G82S.
- 25 30. The method of claim 22 or 23, wherein the subject's cells in which the amount of RAGE G82S is to be reduced are cells which mediate an inflammatory reaction.
- 30 31. The method of claim 22 or 23, wherein the subject is human.
32. The method of claim 22 or 23, wherein the inflammatory disorder is rheumatoid arthritis.

33. Soluble RAGE G82S or a ligand-binding portion thereof.
34. A composition comprising a pharmaceutically acceptable carrier and (i) soluble RAGE G82S, (ii) a ligand-binding portion soluble RAGE G82S, and/or (iii) a nucleic acid which specifically inhibits the expression of RAGE G82S in a cell expressing same.
35. An article of manufacture comprising a packaging material having therein (i) an agent that inhibits binding between RAGE G82S and a ligand thereof, and/or (ii) a nucleic acid which specifically inhibits the expression of RAGE G82S in a cell expressing same, wherein the packaging material has affixed thereto a label indicating a use for the agent and/or nucleic acid for treating an inflammatory disorder in a subject.
36. The article of claim 35, wherein the subject is a human.
37. The article of claim 35, wherein the inflammatory disorder is rheumatoid arthritis and the agent comprises (i) soluble RAGE G82S, (ii) a ligand-binding portion of soluble RAGE G82S, or (iii) an antibody directed to RAGE G82S.
38. The article of claim 37, wherein the agent comprises soluble RAGE G82S.

39. The article of claim 37, wherein the agent comprises a ligand-binding portion of soluble RAGE G82S.

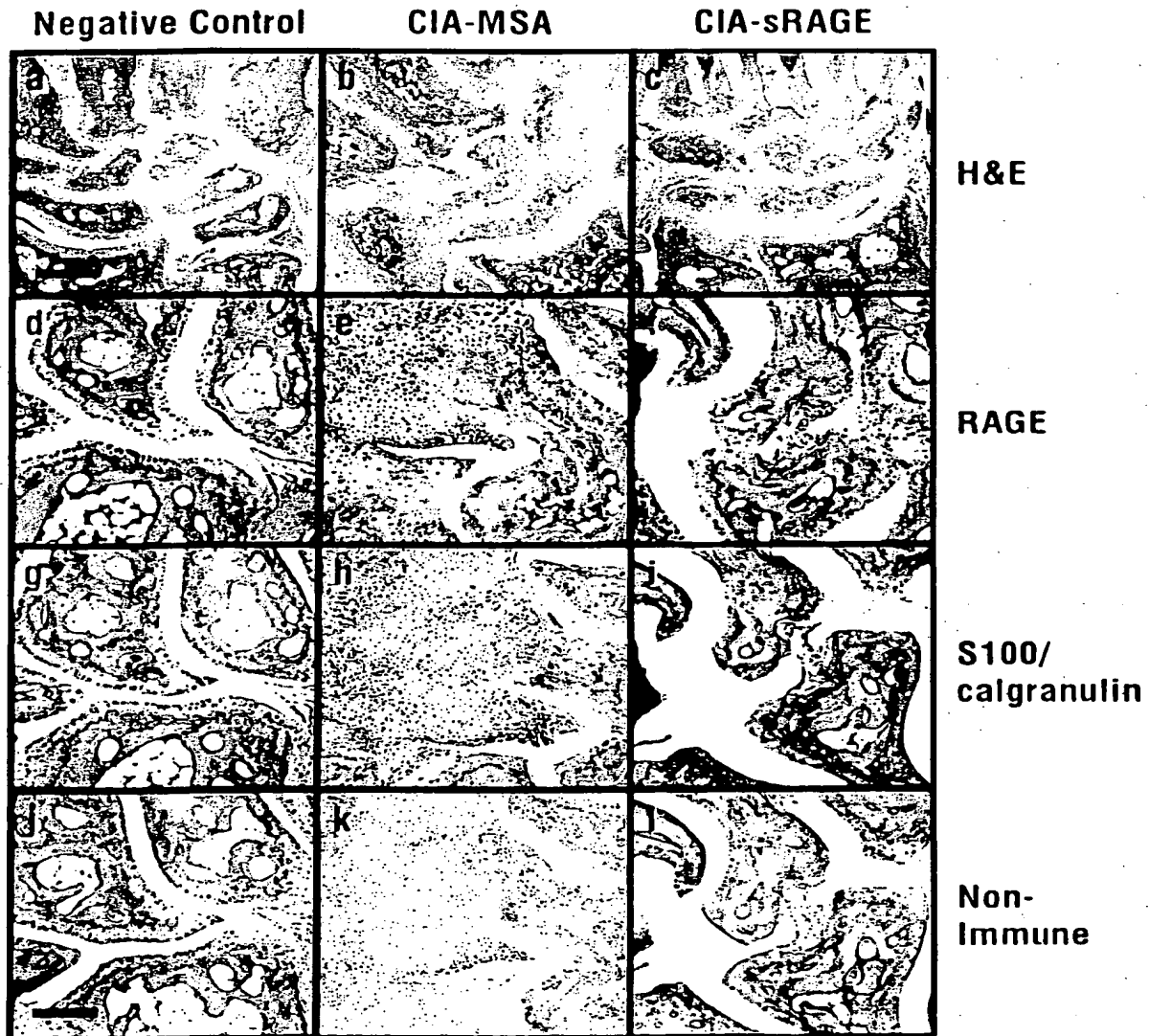
5 40. The article of claim 37, wherein the agent comprises an antibody directed to soluble RAGE G82S.

RAGE G82S-RELATED METHODS AND COMPOSITIONS FOR TREATING
INFLAMMATORY DISORDERS

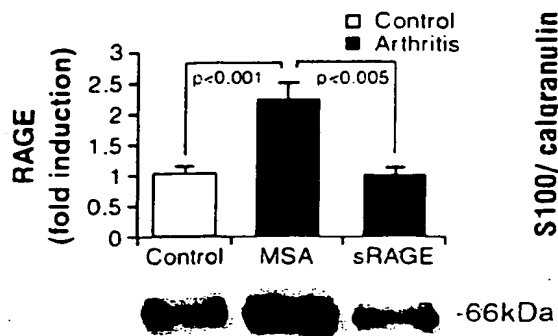
5 Abstract of the Disclosure

This invention provides methods, compositions and
articles of manufacture for inhibiting the onset of and
treating inflammatory disorders such as rheumatoid
10 arthritis. The instant invention is based on the
blockade of RAGE G82S function.

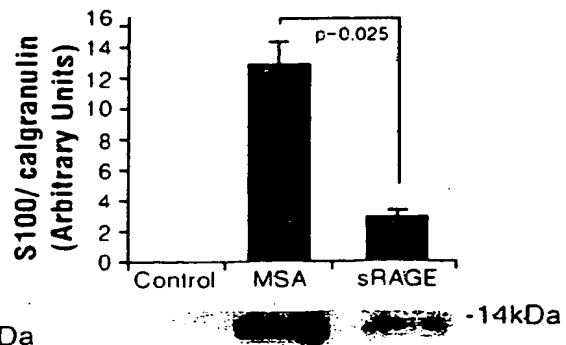
Figures 1A-1N



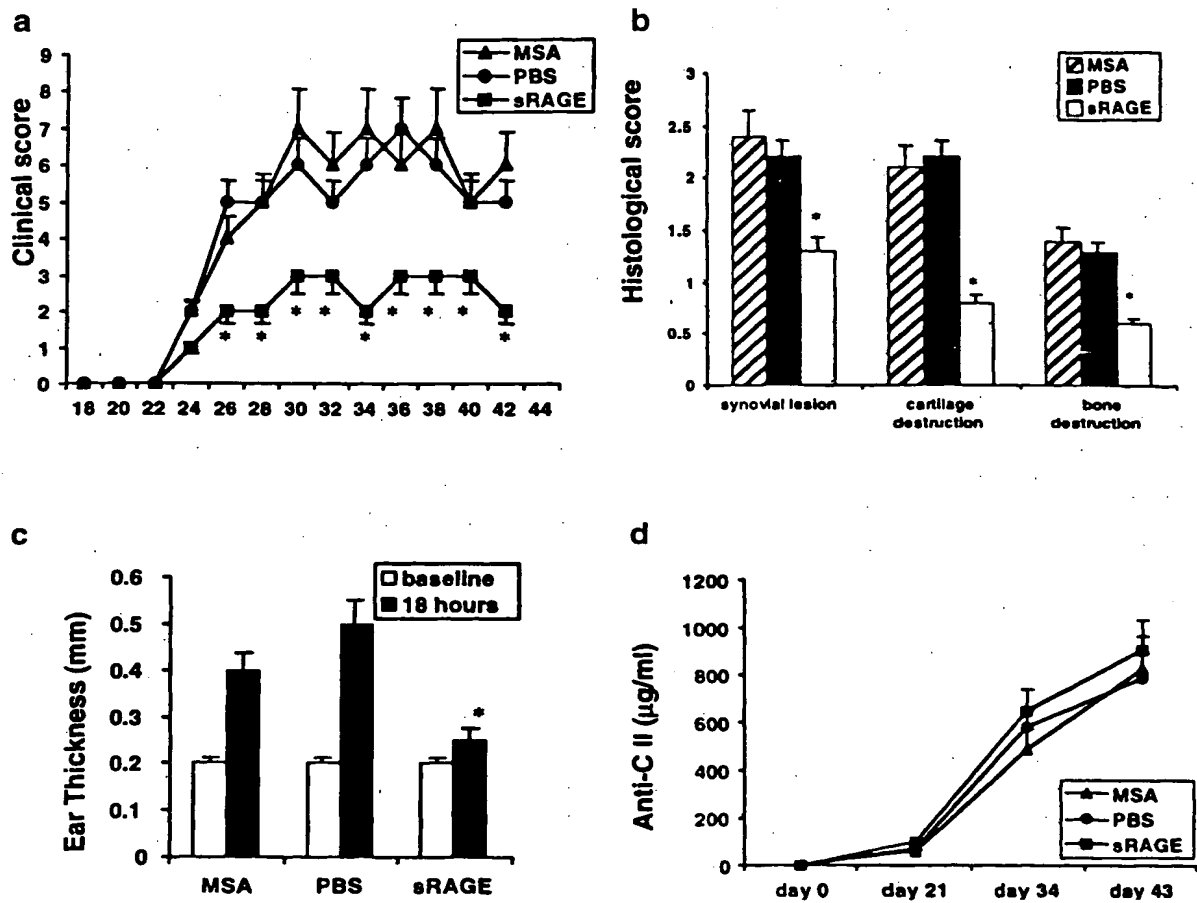
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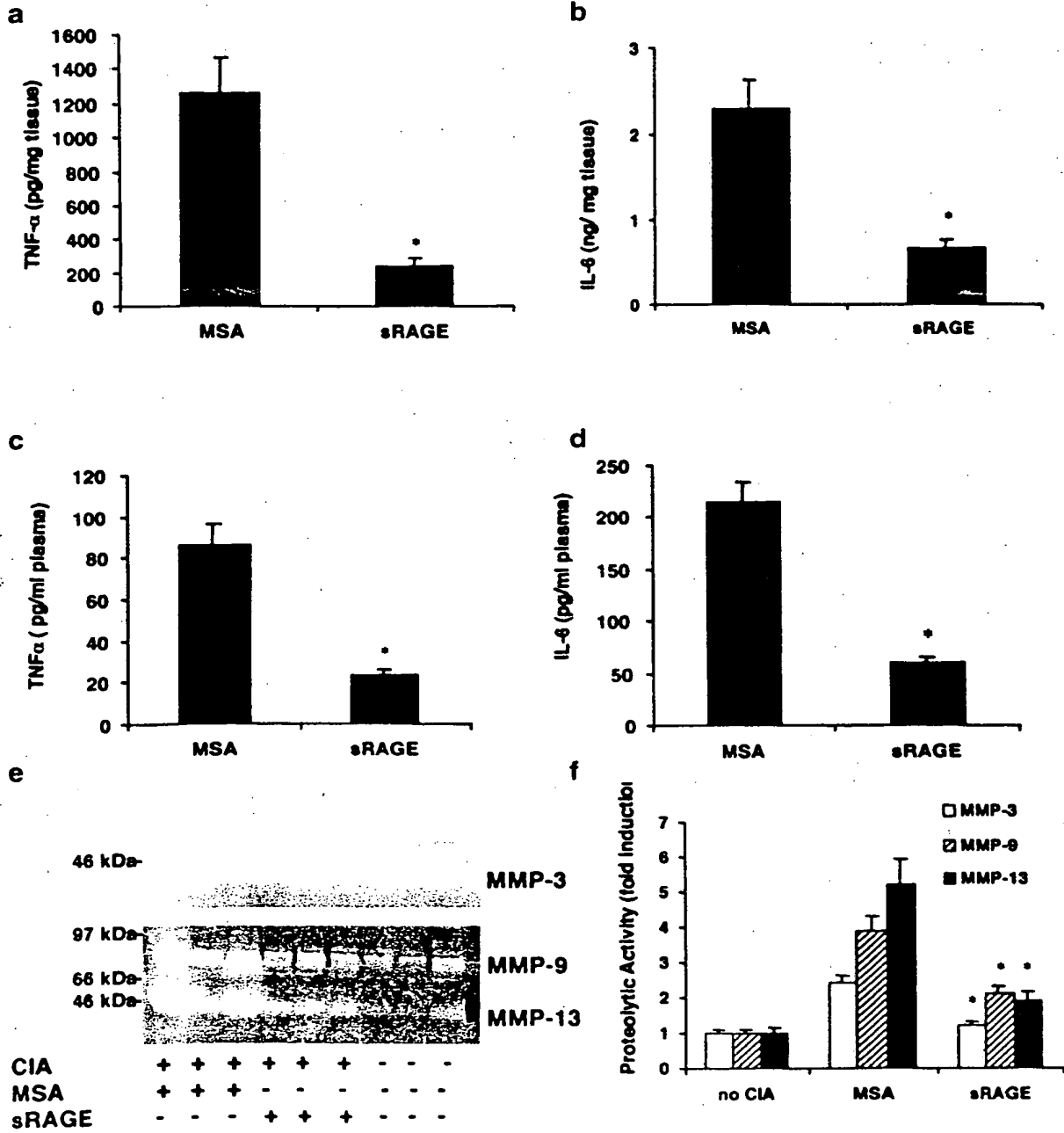
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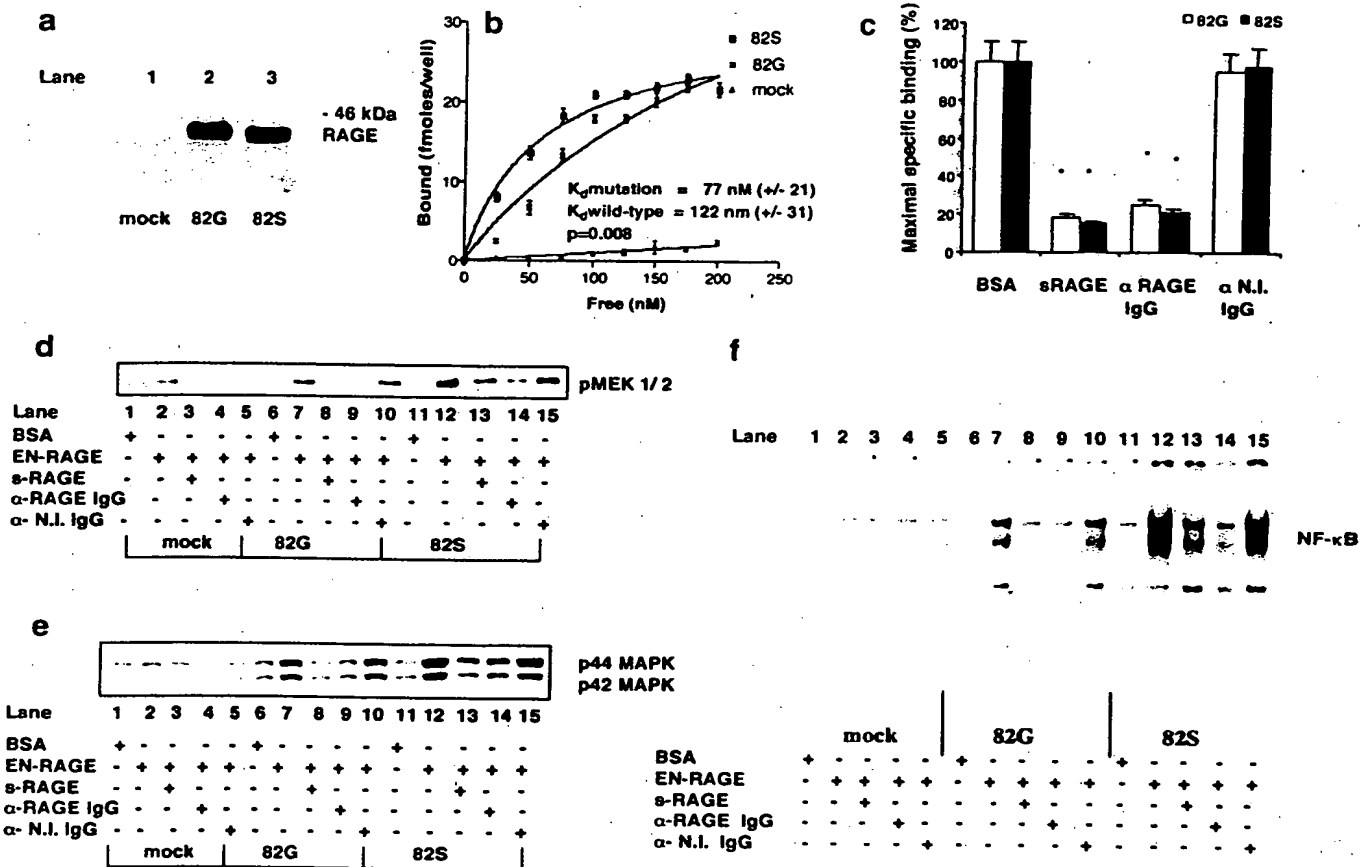
Figures 2A-2D



Figures 3A-3F



Figures 4A-F



Figures 5A-5H

